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(71) Applicant: CORIXA CORPORATION [US/US]; Suit 1124 Columbia Street, Seattle, WA 98104 (US).	te 200	
(72) Inventors: REED, Steven, G.; 2843 - 122nd Place Bellevie, WA 98005 (US). LODES, Michael, J.; 36th Avenue S.W., Seattle, WA 98126 (US). FRUE Tony, N.; P.O. Box 99232, Seattle, WA 99232-0232	9223 DAKIS 2 (US)	
MOHAMATH, Raodoh; 4205 South Morgan, Seattl 98118 (US).	. 172	

(54) Title: COMPOUNDS FOR THERAPY AND DIAGNOSIS OF LUNG CANCER AND METHODS FOR THEIR USE

(57) Abstract

Compounds and methods for treating lung cancer are provided. The inventive compounds include polypeptides containing at least a portion of a lung tumor protein. Vaccines and pharmaceutical compositions for immunotherapy of lung cancer comprising such polypeptides, or polynucleotides encoding such polypeptides, are also provided, together with polynucleotides for preparing the inventive polypeptides.

COMPOUNDS FOR THERAPY AND DIAGNOSIS OF LUNG CANCER AND METHODS FOR THEIR USE

TECHNICAL FIELD

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The present invention relates generally to compositions and methods for the treatment of lung cancer. The invention is more specifically related to nucleotide sequences that are preferentially expressed in lung tumor tissue, together with polypeptides encoded by such nucleotide sequences. The inventive nucleotide sequences and polypeptides may be used in vaccines and pharmaceutical compositions for the treatment of lung cancer.

BACKGROUND OF THE INVENTION

Lung cancer is the primary cause of cancer death among both men and women in the U.S., with an estimated 172,000 new cases being reported in 1994. The five-year survival rate among all lung cancer patients, regardless of the stage of disease at diagnosis, is only 13%. This contrasts with a five-year survival rate of 46% among cases detected while the disease is still localized. However, only 16% of lung cancers are discovered before the disease has spread.

Early detection is difficult since clinical symptoms are often not seen until the disease has reached an advanced stage. Currently, diagnosis is aided by the use of chest x-rays, analysis of the type of cells contained in sputum and fiberoptic examination of the bronchial passages. Treatment regimens are determined by the type and stage of the cancer, and include surgery, radiation therapy and/or chemotherapy. In spite of considerable research into therapies for the disease, lung cancer remains difficult to treat.

Accordingly, there remains a need in the art for improved vaccines, treatment methods and diagnostic techniques for lung cancer.

SUMMARY OF THE INVENTION

Briefly stated, the present invention provides compounds and methods for the therapy of lung cancer. In a first aspect, isolated polynucleotides encoding lung tumor polypeptides are provided, such polynucleotides comprising a nucleotide sequence selected

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herein; and (b) detecting in the sample a protein or polypeptide that binds to the binding agent. In preferred embodiments, the binding agent is an antibody, most preferably a monoclonal antibody.

In related aspects, methods are provided for monitoring the progression of lung cancer in a patient, comprising: (a) contacting a biological sample obtained from a patient with a binding agent that is capable of binding to one of the polypeptides disclosed herein; (b) determining in the sample an amount of a protein or polypeptide that binds to the binding agent; (c) repeating steps (a) and (b); and comparing the amounts of polypeptide detected in steps (b) and (c).

Within related aspects, the present invention provides antibodies, preferably monoclonal antibodies, that bind to the inventive polypeptides, as well as diagnostic kits comprising such antibodies, and methods of using such antibodies to inhibit the development of lung cancer.

The present invention further provides methods for detecting lung cancer comprising: (a) obtaining a biological sample from a patient; (b) contacting the sample with a first and a second oligonucleotide primer in a polymerase chain reaction, at least one of the oligonucleotide primers being specific for a polynucleotide that encodes one of the polypeptides disclosed herein; and (c) detecting in the sample a DNA sequence that amplifies in the presence of the first and second oligonucleotide primers. In a preferred embodiment, at least one of the oligonucleotide primers comprises at least about 10 contiguous nucleotides of a polynucleotide comprising a sequence selected from the group consisting of SEQ ID NO: 1-31, 49-55, 63, 64, 66, 68-72, 78-80, 84-92, 102-110, 116-120 and 126-181.

In a further aspect, the present invention provides a method for detecting lung cancer in a patient comprising: (a) obtaining a biological sample from the patient; (b) contacting the sample with an oligonucleotide probe specific for a polynucleotide that encodes one of the polypeptides disclosed herein; and (c) detecting in the sample a DNA sequence that hybridizes to the oligonucleotide probe. Preferably, the oligonucleotide probe comprises at least about 15 contiguous nucleotides of a polynucleotide comprising a sequence selected from the group consisting of SEQ ID NO: 1-31, 49-55, 63, 64, 66, 68-72, 78-80, 84-92, 102-110, 116-120 and 126-181. In related aspects, diagnostic kits comprising the above oligonucleotide probes or primers are provided.

SEQ ID NO: 14 is the determined cDNA sequence for L355C1.cons SEQ ID NO: 15 is the determined cDNA sequence for L366C1.cons SEQ ID NO: 16 is the determined cDNA sequence for L163C1a SEQ ID NO: 17 is the determined cDNA sequence for LT86-1 SEQ ID NO: 18 is the determined cDNA sequence for LT86-2 SEQ ID NO: 19 is the determined cDNA sequence for LT86-3 Sale State of the SEQ ID NO: 20 is the determined cDNA sequence for LT86-4 organization and dis SEQ ID NO: 21 is the determined cDNA sequence for LT86-5 SEQ ID NO: 22 is the determined cDNA sequence for LT86-6 SEQ ID NO: 23 is the determined cDNA sequence for LT86-7 SEQ ID NO: 24 is the determined cDNA sequence for LT86-8 SEQ ID NO: 25 is the determined cDNA sequence for LT86-9 SEQ ID NO: 26 is the determined cDNA sequence for LT86-10 SEQ ID NO: 27 is the determined cDNA sequence for LT86-11 SEQ ID NO: 28 is the determined cDNA sequence for LT86-12 SEQ ID NO: 29 is the determined cDNA sequence for LT86-13 SEQ ID NO: 30 is the determined cDNA sequence for LT86-14 SEQ ID NO: 31 is the determined cDNA sequence for LT86-15 SEQ ID NO: 32 is the predicted amino acid sequence for LT86-1 , 20 SEQ ID NO: 33 is the predicted amino acid sequence for LT86-2 SEQ ID NO: 34 is the predicted amino acid sequence for LT86-3 SEQ ID NO: 35 is the predicted amino acid sequence for LT86-4 SEQ ID NO: 36 is the predicted amino acid sequence for LT86-5 SEQ ID NO: 37 is the predicted amino acid sequence for LT86-6 SEQ ID NO: 38 is the predicted amino acid sequence for LT86-7 SEQ ID NO: 39 is the predicted amino acid sequence for LT86-8 SEQ ID NO: 40 is the predicted amino acid sequence for LT86-9 SEQ ID NO: 41 is the predicted amino acid sequence for LT86-10 SEQ ID NO: 42 is the predicted amino acid sequence for LT86-11 SEQ ID NO: 43 is the predicted amino acid sequence for LT86-12

SEQ ID NO: 74 is the predicted amino acid sequence for LT86-21 SEQ ID NO: 75 is the predicted amino acid sequence for LT86-22 SEQ ID NO: 76 is the predicted amino acid sequence for LT86-26 SEQ ID NO: 77 is the predicted amino acid sequence for LT86-27 SEQ ID NO: 78 is the determined extended cDNA sequence for L86S-12 SEQ ID NO: 79 is the determined extended cDNA sequence for L86S-36 SEQ ID NO: 80 is the determined extended cDNA sequence for L86S-46 SEQ ID NO: 81 is the predicted extended amino acid sequence for L86S-12 SEQ ID NO: 82 is the predicted extended amino acid sequence for L86S-36 SEQ ID NO: 83 is the predicted extended amino acid sequence for L86S-46 SEQ ID NO: 84 is the determined 5'cDNA sequence for L86S-6 SEQ ID NO: 85 is the determined 5'cDNA sequence for L86S-11 SEO ID NO: 86 is the determined 5'cDNA sequence for L86S-14 SEQ ID NO: 87 is the determined 5'cDNA sequence for L86S-29 SEQ ID NO: 88 is the determined 5'cDNA sequence for L86S-34 15 SEQ ID NO: 89 is the determined 5'cDNA sequence for L86S-39 SEQ ID NO: 90 is the determined 5'cDNA sequence for L86S-47 SEQ ID NO: 91 is the determined 5'cDNA sequence for L86S-49 SEQ ID NO: 92 is the determined 5'cDNA sequence for L86S-51 SEQ ID NO: 93 is the predicted amino acid sequence for L86S-6 20 SEQ ID NO: 94 is the predicted amino acid sequence for L86S-11 SEQ ID NO: 95 is the predicted amino acid sequence for L86S-14 SEQ ID NO: 96 is the predicted amino acid sequence for L86S-29 SEQ ID NO: 97 is the predicted amino acid sequence for L86S-34 25 SEQ ID NO: 98 is the predicted amino acid sequence for L86S-39 SEQ ID NO: 99 is the predicted amino acid sequence for L86S-47. SEQ ID NO: 100 is the predicted amino acid sequence for L86S-49 SEQ ID NO: 101 is the predicted amino acid sequence for L86S-51 SEQ ID NO: 102 is the determined DNA sequence for SLT-T1

SEQ ID NO: 103 is the determined 5' cDNA sequence for SLT-T2

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SEQ ID NO: 134 is the determined cDNA sequence for PSLT-69 SEQ ID NO: 135 is the determined cDNA sequence for PSLT-71 SEQ ID NO: 136 is the determined cDNA sequence for PSLT-73 SEQ ID NO: 137 is the determined cDNA sequence for PSLT-79 SEQ ID NO: 138 is the determined cDNA sequence for PSLT-03 SEQ ID NO: 139 is the determined cDNA sequence for PSLT-09 SEQ ID NO: 140 is the determined cDNA sequence for PSLT-011 SEQ ID NO: 141 is the determined cDNA sequence for PSLT-041 SEQ ID NO: 142 is the determined cDNA sequence for PSLT-62 SEQ ID NO: 143 is the determined cDNA sequence for PSLT-6 SEQ ID NO: 144 is the determined cDNA sequence for PSLT-37 SEQ ID NO: 145 is the determined cDNA sequence for PSLT-74 SEQ ID NO: 146 is the determined cDNA sequence for PSLT-010 SEQ ID NO: 147 is the determined cDNA sequence for PSLT-012 SEQ ID NO: 148 is the determined cDNA sequence for PSLT-037 SEQ ID NO: 149 is the determined 5' cDNA sequence for SAL-3 SEQ ID NO: 150 is the determined 5' cDNA sequence for SAL-24 SEQ ID NO: 151 is the determined 5' cDNA sequence for SAL-25 SEQ ID NO: 152 is the determined 5' cDNA sequence for SAL-33 SEQ ID NO: 153 is the determined 5' cDNA sequence for SAL-50 SEQ ID NO: 154 is the determined 5' cDNA sequence for SAL-57 SEQ ID NO: 155 is the determined 5' cDNA sequence for SAL-66 SEQ ID NO: 156 is the determined 5' cDNA sequence for SAL-82 SEQ ID NO: 157 is the determined 5' cDNA sequence for SAL-99 SEQ ID NO: 158 is the determined 5' cDNA sequence for SAL-104 25 SEQ ID NO: 159 is the determined 5' cDNA sequence for SAL-109 SEQ ID NO: 160 is the determined 5' cDNA sequence for SAL-5 SEQ ID NO: 161 is the determined 5° cDNA sequence for SAL-8 SEQ ID NO: 162 is the determined 5' cDNA sequence for SAL-12 SEQ ID NO: 163 is the determined 5' cDNA sequence for SAL-14

SEQ ID NO: 194 is the predicted amino acid sequence for SAL-5 SEQ ID NO: 195 is the predicted amino acid sequence for SAL-8 SEQ ID NO: 196 is the predicted amino acid sequence for SAL-12 SEQ ID NO: 197 is the predicted amino acid sequence for SAL-14 5 SEQ ID NO: 198 is the predicted amino acid sequence for SAL-16 SEQ ID NO: 199 is the predicted amino acid sequence for SAL-23 SEQ ID NO: 200 is the predicted amino acid sequence for SAL-26 SEQ ID NO: 201 is the predicted amino acid sequence for SAL-29 SEQ ID NO: 202 is the predicted amino acid sequence for SAL-32 SEQ ID NO: 203 is the predicted amino acid sequence for SAL-39 SEQ ID NO: 204 is the predicted amino acid sequence for SAL-42 SEQ ID NO: 205 is the predicted amino acid sequence for SAL-43 SEQ ID NO: 206 is the predicted amino acid sequence for SAL-44 SEQ ID NO: 207 is the predicted amino acid sequence for SAL-48 15 SEQ ID NO: 208 is the predicted amino acid sequence for SAL-68 SEQ ID NO: 209 is the predicted amino acid sequence for SAL-72 SEQ ID NO: 210 is the predicted amino acid sequence for SAL-77 SEQ ID NO: 211 is the predicted amino acid sequence for SAL-86 SEQ ID NO: 212 is the predicted amino acid sequence for SAL-88 gare of the colony SEQ ID NO: 213 is the predicted amino acid sequence for SAL-93 20 SEQ ID NO: 214 is the predicted amino acid sequence for SAL-100 SEQ ID NO: 215 is the predicted amino acid sequence for SAL-105 SEQ ID NO: 216 is a second predicted amino acid sequence for SAL-50 spread from the

25 DETAILED DESCRIPTION OF THE INVENTION

As noted above, the present invention is generally directed to compositions and methods for the therapy of lung cancer. The compositions described herein include polypeptides, fusion proteins and polynucleotides. Also included within the present invention are molecules (such as an antibody or fragment thereof) that bind to the inventive polypeptides. Such molecules are referred to herein as "binding agents."

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of the proteins described herein may be identified in antibody binding assays. Such assays may generally be performed using any of a variety of means known to those of ordinary skill in the art, as described, for example, in Harlow and Lane, Antibodies: A Laboratory Manual, Cold Spring Harbor Laboratory, Cold Spring Harbor, NY, 1988. For example, a polypeptide may be immobilized on a solid support (as described below) and contacted with patient sera to allow binding of antibodies within the sera to the immobilized polypeptide. Unbound sera may then be removed and bound antibodies detected using, for example, ¹²⁵I-labeled Protein A. Alternatively, a polypeptide may be used to generate monoclonal and polyclonal antibodies for use in detection of the polypeptide in blood or other fluids of lung cancer patients. Methods for preparing and identifying immunogenic portions of antigens of known sequence are well known in the art and include those summarized in Paul, Fundamental Immunology, 3rd ed., Raven Press, 1993, pp. 243-247.

The term "polynucleotide(s)," as used herein, means a single or double-stranded polymer of deoxyribonucleotide or ribonucleotide bases and includes DNA and corresponding RNA molecules, including HnRNA and mRNA molecules, both sense and anti-sense strands, and comprehends cDNA, genomic DNA and recombinant DNA, as well as wholly or partially synthesized polynucleotides. An HnRNA molecule contains introns and corresponds to a DNA molecule in a generally one-to-one manner. An mRNA molecule corresponds to an HnRNA and DNA molecule from which the introns have been excised. A polynucleotide may consist of an entire gene, or any portion thereof. Operable anti-sense polynucleotides may comprise a fragment of the corresponding polynucleotide, and the definition of "polynucleotide" therefore includes all such operable anti-sense fragments.

The compositions and methods of the present invention also encompass variants of the above polypeptides and polynucleotides.

A polypeptide "variant," as used herein, is a polypeptide that differs from the recited polypeptide only in conservative substitutions and/or modifications, such that the antigenic properties of the polypeptide are retained. In a preferred embodiment, variant polypeptides differ from an identified sequence by substitution, deletion or addition of five amino acids or fewer. Such variants may generally be identified by modifying one of the above polypeptide sequences, and evaluating the antigenic properties of the modified polypeptide using, for example, the representative procedures described herein. Polypeptide

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SSC, 0.5% SDS, 1.0 mM EDTA (pH 8.0); hybridizing at 50°C-65°C, 5X SSC, overnight or, in the event of cross-species homology, at 45°C with 0.5X SSC; followed by washing twice at 65°C for 20 minutes with each of 2X, 0.5X and 0.2X SSC containing 0.1% SDS. Such hybridizing DNA sequences are also within the scope of this invention, as are nucleotide sequences that, due to code degeneracy, encode an immunogenic polypeptide that is encoded by a hybridizing DNA sequence.

Two nucleotide or polypeptide sequences are said to be "identical" if the sequence of nucleotides or amino acid residues in the two sequences is the same when aligned for maximum correspondence as described below. Comparisons between two sequences are typically performed by comparing the sequences over a comparison window to identify and compare local regions of sequence similarity. A "comparison window" as used herein, refers to a segment of at least about 20 contiguous positions, usually 30 to about 75, 40 to about 50, in which a sequence may be compared to a reference sequence of the same number of contiguous positions after the two sequences are optimally aligned.

Optimal alignment of sequences for comparison may be conducted using the Megalign program in the Lasergene suite of bioinformatics software (DNASTAR, Inc., Madison, WI), using default parameters. This program embodies several alignment schemes described in the following references: Dayhoff, M.O. (1978) A model of evolutionary change in proteins - Matrices for detecting distant relationships. In Dayhoff, M.O. (ed.) Atlas of Protein Sequence and Structure, National Biomedical Resarch Foundation, Washington DC Vol. 5, Suppl. 3, pp. 345-358; Hein J. (1990) Unified Approach to Alignment and Phylogenes pp. 626-645 Methods in Enzymology vol. 183; Academic Press, Inc., San Diego, CA; Higgins, D.G. and Sharp, P.M. (1989) Fast and sensitive multiple sequence alignments on a microcomputer CABIOS 5:151-153; Myers, E.W. and Muller W. (1988) Optimal alignments in linear space CABIOS 4:11-17; Robinson, E.D. (1971) Comb. Theor 11:105; Santou, N. Nes, M. (1987) The neighbor joining method. A new method for reconstructing phylogenetic trees Mol. Biol. Evol. 4:406-425; Sneath, P.H.A. and Sokal, R.R. (1973) Numerical Taxonomy - the Principles and Practice of Numerical Taxonomy. Freeman Press, San Francisco, CA; Wilbur, W.J. and Lipman, D.J. (1983) Rapid similarity searches of nucleic acid and protein data banks Proc. Natl. Acad., Sci. USA 80:726-730.

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libraries prepared from SCID mice with mouse anti-tumor sera, as described below in Example 4. Examples of cDNA sequences that may be isolated using this technique are provided in SEQ ID NO: 149-181.

A gene encoding a polypeptide described herein (or a portion thereof) may, alternatively, be amplified from human genomic DNA, or from lung tumor cDNA, via polymerase chain reaction. For this approach, sequence-specific primers may be designed based on the nucleotide sequences provided herein and may be purchased or synthesized. An amplified portion of a specific nucleotide sequence may then be used to isolate the full length gene from a human genomic DNA library or from a lung tumor cDNA library, using well known techniques, such as those described in Sambrook et al., *Molecular Cloning: A Laboratory Manual*, Cold Spring Harbor Laboratories, Cold Spring Harbor, NY (1989).

Once a DNA sequence encoding a polypeptide is obtained, the polypeptide may be produced recombinantly by inserting the DNA sequence into an expression vector and expressing the polypeptide in an appropriate host. Any of a variety of expression vectors known to those of ordinary skill in the art may be employed to express recombinant polypeptides of this invention. Expression may be achieved in any appropriate host cell that has been transformed or transfected with an expression vector containing a polynucleotide that encodes the recombinant polypeptide. Suitable host cells include prokaryotes, yeast and higher eukaryotic cells. Preferably, the host cells employed are *E. coli*, yeast or a mammalian cell line, such as COS or CHO cells. The DNA sequences expressed in this manner may encode naturally occurring polypeptides, portions of naturally occurring polypeptides, or other variants thereof. Supernatants from suitable host/vector systems which secrete the recombinant polypeptide may be first concentrated using a commercially available filter. The concentrate may then be applied to a suitable purification matrix, such as an affinity matrix or ion exchange resin. Finally, one or more reverse phase HPLC steps can be employed to further purify the recombinant polypeptide.

Such techniques may also be used to prepare polypeptides comprising portions or variants of the native polypeptides. Portions and other variants having fewer than about 100 amino acids, and generally fewer than about 50 amino acids, may be generated using techniques well known to those of ordinary skill in the art. For example, such polypeptides may be synthesized using any of the commercially available solid-phase techniques, such as

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extended conformation; (2) their inability to adopt a secondary structure that could interact with functional epitopes on the first and second polypeptides; and (3) the lack of hydrophobic or charged residues that might react with the polypeptide functional epitopes. Preferred peptide linker sequences contain Gly, Asn and Ser residues. Other near neutral amino acids, such as Thr and Ala may also be used in the linker sequence. Amino acid sequences which may be usefully employed as linkers include those disclosed in Maratea et al., Gene 40:39-46, 1985; Murphy et al., Proc. Natl. Acad. Sci. USA 83:8258-8262, 1986; U.S. Patent No. 4,935,233 and U.S. Patent No. 4,751,180. The linker sequence may be from 1 to about 50 amino acids in length. Peptide sequences are not required when the first and second polypeptides have non-essential N-terminal amino acid regions that can be used to separate the functional domains and prevent steric interference.

The ligated DNA sequences are operably linked to suitable transcriptional or translational regulatory elements. The regulatory elements responsible for expression of DNA are located only 5' to the DNA sequence encoding the first polypeptides. Similarly, stop codons require to end translation and transcription termination signals are only present 3' to the DNA sequence encoding the second polypeptide.

Fusion proteins are also provided that comprise a polypeptide of the present invention together with an unrelated immunogenic protein. Preferably the immunogenic protein is capable of eliciting a recall response. Examples of such proteins include tetanus, tuberculosis and hepatitis proteins (see, for example, Stoute et al. New Engl. J. Med., 336:86-91 (1997)).

Polypeptides that comprise an immunogenic portion of a lung tumor protein may generally be used for therapy of lung cancer, wherein the polypeptide stimulates the patient's own immune response to lung tumor cells. The present invention thus provides methods for using one or more of the compounds described herein (which may be polypeptides, polynucleotides or fusion proteins) for immunotherapy of lung cancer in a patient. As used herein, a "patient" refers to any warm-blooded animal, preferably a human. A patient may be afflicted with disease, or may be free of detectable disease. Accordingly, the compounds disclosed herein may be used to treat lung cancer or to inhibit the development of lung cancer. In a preferred embodiment, the compounds are administered

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ordinary skill in the art. The DNA may also be "naked," as described, for example, in published PCT application WO 90/11092, and Ulmer et al., *Science 259*:1745-1749, 1993, reviewed by Cohen, *Science 259*:1691-1692, 1993. The uptake of naked DNA may be increased by coating the DNA onto biodegradable beads, which are efficiently transported into the cells.

Routes and frequency of administration, as well as dosage, will vary from individual to individual and may parallel those currently being used in immunotherapy of other diseases. In general, the pharmaceutical compositions and vaccines may be administered by injection (e.g., intracutaneous, intramuscular, intravenous or subcutaneous), intranasally (e.g., by aspiration) or orally. Between 1 and 10 doses may be administered over a 3-24 week period. Preferably, 4 doses are administered, at an interval of 3 months, and booster administrations may be given periodically thereafter. Alternate protocols may be appropriate for individual patients. A suitable dose is an amount of polypeptide or DNA that is effective to raise an immune response (cellular and/or humoral) against lung tumor cells in a treated patient. A suitable immune response is at least 10-50% above the basal (i.e., untreated) level. In general, the amount of polypeptide present in a dose (or produced in situ by the DNA in a dose) ranges from about 1 pg to about 100 mg per kg of host, typically from about 10 pg to about 1 mg, and preferably from about 100 pg to about 1 µg. Suitable dose sizes will vary with the size of the patient, but will typically range from about 0.01 mL to about 5 mL.

While any suitable carrier known to those of ordinary skill in the art may be employed in the pharmaceutical compositions of this invention, the type of carrier will vary depending on the mode of administration. For parenteral administration, such as subcutaneous injection, the carrier preferably comprises water, saline, alcohol, a lipid, a wax and/or a buffer. For oral administration, any of the above carriers or a solid carrier, such as mannitol, lactose, starch, magnesium stearate, sodium saccharine, talcum, cellulose, glucose, sucrose, and/or magnesium carbonate, may be employed. Biodegradable microspheres (e.g., polylactic glycolide) may also be employed as carriers for the pharmaceutical compositions of this invention. Suitable biodegradable microspheres are disclosed, for example, in U.S. Patent Nos. 4.897,268 and 5,075,109.

(Natural Killer cells, lymphokine-activated killer cells), B cells, or antigen presenting cells (such as dendritic cells and macrophages) expressing the disclosed antigens. The polypeptides disclosed herein may also be used to generate antibodies or anti-idiotypic antibodies (as in U.S. Patent No. 4,918,164), for passive immunotherapy.

The predominant method of procuring adequate numbers of T-cells for adoptive immunotherapy is to grow immune T-cells in vitro. Culture conditions for expanding single antigen-specific T-cells to several billion in number with retention of antigen recognition in vivo are well known in the art. These in vitro culture conditions typically utilize intermittent stimulation with antigen, often in the presence of cytokines, such as IL-2, and non-dividing feeder cells. As noted above, the immunoreactive polypeptides described herein may be used to rapidly expand antigen-specific T cell cultures in order to generate sufficient number of cells for immunotherapy. In particular, antigen-presenting cells, such as dendritic, macrophage or B-cells, may be pulsed with immunoreactive polypeptides or transfected with a polynucleotide sequence(s), using standard techniques well known in the art. For cultured T-cells to be effective in therapy, the cultured T-cells must be able to grow and distribute widely and to survive long term in vivo. Studies have demonstrated that cultured T-cells can be induced to grow in vivo and to survive long term in substantial numbers by repeated stimulation with antigen supplemented with IL-2 (see, for example, Cheever et al. *Ibid*).

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The polypeptides disclosed herein may also be employed to generate and/or isolate tumor-reactive T-cells, which can then be administered to the patient. In one technique, antigen-specific T-cell lines may be generated by *in vivo* immunization with short peptides corresponding to immunogenic portions of the disclosed polypeptides. The resulting antigen specific CD8+ CTL clones may be isolated from the patient, expanded using standard tissue culture techniques, and returned to the patient.

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Alternatively, peptides corresponding to immunogenic portions of the polypeptides may be employed to generate tumor reactive T cell subsets by selective in vitro stimulation and expansion of autologous T cells to provide antigen-specific T cells which may be subsequently transferred to the patient as described, for example, by Chang et al. (Crit. Rev. Oncol. Hematol., 22(3), 213, 1996).

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at least about 80%, and preferably at least about 90%) of the patients for which lung cancer would be indicated using the full length protein, and that indicate the absence of lung cancer in substantially all of those samples that would be negative when tested with full length protein. The representative assays described below, such as the two-antibody sandwich assay, may generally be employed for evaluating the ability of a binding agent to detect metastatic human lung tumors.

The ability of a polypeptide prepared as described herein to generate antibodies capable of detecting primary or metastatic human lung tumors may generally be evaluated by raising one or more antibodies against the polypeptide (using, for example, a representative method described herein) and determining the ability of such antibodies to detect such tumors in patients. This determination may be made by assaying biological samples from patients with and without primary or metastatic lung cancer for the presence of a polypeptide that binds to the generated antibodies. Such test assays may be performed, for example, using a representative procedure described below. Polypeptides that generate 15 antibodies capable of detecting at least 20% of primary or metastatic lung tumors by such procedures are considered to be useful in assays for detecting primary or metastatic human lung tumors. Polypeptide specific antibodies may be used alone or in combination to improve sensitivity.

Polypeptides capable of detecting primary or metastatic human lung tumors 20 may be used as markers for diagnosing lung cancer or for monitoring disease progression in patients. In one embodiment, lung cancer in a patient may be diagnosed by evaluating a biological sample obtained from the patient for the level of one or more of the above polypeptides, relative to a predetermined cut-off value. As used herein, suitable "biological samples" include blood, sera, urine and/or lung secretions.

The level of one or more of the above polypeptides may be evaluated using any binding agent specific for the polypeptide(s). A "binding agent," in the context of this invention, is any agent (such as a compound or a cell) that binds to a polypeptide as described above. As used herein, "binding" refers to a noncovalent association between two separate molecules (each of which may be free (i.e., in solution) or present on the surface of a cell or a solid support), such that a "complex" is formed. Such a complex may be free or immobilized (either covalently or noncovalently) on a support material. The ability to bind may generally

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be immobilized on the solid support using a variety of techniques known to those of skill in the art, which are amply described in the patent and scientific literature. In the context of the present invention, the term "immobilization" refers to both noncovalent association, such as adsorption, and covalent attachment (which may be a direct linkage between the antigen and functional groups on the support or may be a linkage by way of a cross-linking agent). Immobilization by adsorption to a well in a microtiter plate or to a membrane is preferred. In such cases, adsorption may be achieved by contacting the binding agent, in a suitable buffer, with the solid support for a suitable amount of time. The contact time varies with temperature, but is typically between about 1 hour and about 1 day. In general, contacting a well of a plastic microtiter plate (such as polystyrene or polyvinylchloride) with an amount of binding agent ranging from about 10 ng to about 10 µg, and preferably about 100 ng to about 1 µg, is sufficient to immobilize an adequate amount of binding agent.

Covalent attachment of binding agent to a solid support may generally be achieved by first reacting the support with a bifunctional reagent that will react with both the support and a functional group, such as a hydroxyl or amino group, on the binding agent. For example, the binding agent may be covalently attached to supports having an appropriate polymer coating using benzoquinone or by condensation of an aldehyde group on the support with an amine and an active hydrogen on the binding partner (see, e.g., Pierce Immunotechnology Catalog and Handbook, 1991, at A12-A13).

In certain embodiments, the assay is a two-antibody sandwich assay. This assay may be performed by first contacting an antibody that has been immobilized on a solid support, commonly the well of a microtiter plate, with the sample, such that polypeptides within the sample are allowed to bind to the immobilized antibody. Unbound sample is then removed from the immobilized polypeptide-antibody complexes and a second antibody (containing a reporter group) capable of binding to a different site on the polypeptide is added. The amount of second antibody that remains bound to the solid support is then determined using a method appropriate for the specific reporter group.

More specifically, once the antibody is immobilized on the support as described above, the remaining protein binding sites on the support are typically blocked. Any suitable blocking agent known to those of ordinary skill in the art, such as bovine serum albumin or Tween 20TM (Sigma Chemical Co., St. Louis, MO). The immobilized antibody is

that corresponds to a predetermined cut-off value. In one preferred embodiment, the cut-off value is the average mean signal obtained when the immobilized antibody is incubated with samples from patients without lung cancer. In general, a sample generating a signal that is three standard deviations above the predetermined cut-off value is considered positive for lung cancer. In an alternate preferred embodiment, the cut-off value is determined using a Receiver Operator Curve, according to the method of Sackett et al., Clinical Epidemiology: A Basic Science for Clinical Medicine, Little Brown and Co., 1985, p. 106-7. Briefly, in this embodiment, the cut-off value may be determined from a plot of pairs of true positive rates (i.e., sensitivity) and false positive rates (100%-specificity) that correspond to each possible cut-off value for the diagnostic test result. The cut-off value on the plot that is the closest to the upper left-hand corner (i.e., the value that encloses the largest area) is the most accurate cut-off value, and a sample generating a signal that is higher than the cut-off value determined by this method may be considered positive. Alternatively, the cut-off value may be shifted to the left along the plot, to minimize the false positive rate, or to the right, to minimize the false negative rate. In general, a sample generating a signal that is higher than the cut-off value determined by this method is considered positive for lung cancer.

In a related embodiment, the assay is performed in a flow-through or strip test format, wherein the antibody is immobilized on a membrane, such as nitrocellulose. In the flow-through test, polypeptides within the sample bind to the immobilized antibody as the sample passes through the membrane. A second, labeled antibody then binds to the antibody-polypeptide complex as a solution containing the second antibody flows through the membrane. The detection of bound second antibody may then be performed as described above. In the strip test format, one end of the membrane to which antibody is bound is immersed in a solution containing the sample. The sample migrates along the membrane through a region containing second antibody and to the area of immobilized antibody. Concentration of second antibody at the area of immobilized antibody indicates the presence of lung cancer. Typically, the concentration of second antibody at that site generates a pattern, such as a line, that can be read visually. The absence of such a pattern indicates a negative result. In general, the amount of antibody immobilized on the membrane is selected to generate a visually discernible pattern when the biological sample contains a level of polypeptide that would be sufficient to generate a positive signal in the two-antibody

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of immortal cell lines capable of producing antibodies having the desired specificity (i.e., reactivity with the polypeptide of interest). Such cell lines may be produced, for example, from spleen cells obtained from an animal immunized as described above. The spleen cells are then immortalized by, for example, fusion with a myeloma cell fusion partner, preferably one that is syngeneic with the immunized animal. A variety of fusion techniques may be employed. For example, the spleen cells and myeloma cells may be combined with a nonionic detergent for a few minutes and then plated at low density on a selective medium that supports the growth of hybrid cells, but not myeloma cells. A preferred selection technique uses HAT (hypoxanthine, aminopterin, thymidine) selection. After a sufficient time, usually about 1 to 2 weeks, colonies of hybrids are observed. Single colonies are selected and tested for binding activity against the polypeptide. Hybridomas having high reactivity and specificity are preferred.

Monoclonal antibodies may be isolated from the supernatants of growing hybridoma colonies. In addition, various techniques may be employed to enhance the yield, such as injection of the hybridoma cell line into the peritoneal cavity of a suitable vertebrate host, such as a mouse. Monoclonal antibodies may then be harvested from the ascites fluid or the blood. Contaminants may be removed from the antibodies by conventional techniques, such as chromatography, gel filtration, precipitation, and extraction. The polypeptides of this invention may be used in the purification process in, for example, an affinity chromatography step.

Monoclonal antibodies of the present invention may also be used as therapeutic reagents, to diminish or eliminate lung tumors. The antibodies may be used on their own (for instance, to inhibit metastases) or coupled to one or more therapeutic agents. Suitable agents in this regard include radionuclides, differentiation inducers, drugs, toxins, and derivatives thereof. Preferred radionuclides include ⁹⁰Y, ¹²³I, ¹²⁵I, ¹³¹I, ¹⁸⁶Re, ¹⁸⁸Re, ²¹¹At, and ²¹²Bi. Preferred drugs include methotrexate, and pyrimidine and purine analogs. Preferred differentiation inducers include phorbol esters and butyric acid. Preferred toxins include ricin, abrin, diptheria toxin, cholera toxin, gelonin, Pseudomonas exotoxin, Shigella toxin, and pokeweed antiviral protein.

A therapeutic agent may be coupled (e.g., covalently bonded) to a suitable monoclonal antibody either directly or indirectly (e.g., via a linker group). A direct reaction

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be prepared in a variety of ways. For example, more than one agent may be coupled directly to an antibody molecule, or linkers which provide multiple sites for attachment can be used. Alternatively, a carrier can be used.

A carrier may bear the agents in a variety of ways, including covalent bonding either directly or via a linker group. Suitable carriers include proteins such as albumins (e.g., U.S. Patent No. 4,507,234, to Kato et al.), peptides and polysaccharides such as aminodextran (e.g., U.S. Patent No. 4,699,784, to Shih et al.). A carrier may also bear an agent by noncovalent bonding or by encapsulation, such as within a liposome vesicle (e.g., U.S. Patent Nos. 4,429,008 and 4,873,088). Carriers specific for radionuclide agents include radiohalogenated small molecules and chelating compounds. For example, U.S. Patent No. 4,735,792 discloses representative radiohalogenated small molecules and their synthesis. A radionuclide chelate may be formed from chelating compounds that include those containing nitrogen and sulfur atoms as the donor atoms for binding the metal, or metal oxide, radionuclide. For example, U.S. Patent No. 4,673,562, to Davison et al. discloses representative chelating compounds and their synthesis.

A variety of routes of administration for the antibodies and immunoconjugates may be used. Typically, administration will be intravenous, intramuscular, subcutaneous or in the bed of a resected tumor. It will be evident that the precise dose of the antibody/immunoconjugate will vary depending upon the antibody used, the antigen density on the tumor, and the rate of clearance of the antibody.

Diagnostic reagents of the present invention may also comprise DNA sequences encoding one or more of the above polypeptides, or one or more portions thereof. For example, at least two oligonucleotide primers may be employed in a polymerase chain reaction (PCR) based assay to amplify lung tumor-specific cDNA derived from a biological sample, wherein at least one of the oligonucleotide primers is specific for a polynucleotide encoding a lung tumor protein of the present invention. The presence of the amplified cDNA is then detected using techniques well known in the art, such as gel electrophoresis. Similarly, oligonucleotide probes specific for a polynucleotide encoding a lung tumor protein of the present invention may be used in a hybridization assay to detect the presence of an inventive polypeptide in a biological sample.

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The following Examples are offered by way of illustration and not by way of limitation.

EXAMPLES

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PREPARATION OF LUNG TUMOR-SPECIFIC CDNA SEQUENCES USING DIFFERENTIAL DISPLAY RT-PCR

This example illustrates the preparation of cDNA molecules encoding lung tumor-specific polypeptides using a differential display screen.

Tissue samples were prepared from breast tumor and normal tissue of a patient with lung cancer that was confirmed by pathology after removal of samples from the patient. Normal RNA and tumor RNA was extracted from the samples and mRNA was isolated and converted into cDNA using a (dT)₁₂AG (SEQ ID NO: 47) anchored 3' primer. Differential display PCR was then executed using a randomly chosen primer (SEQ ID NO: 48). Amplification conditions were standard buffer containing 1.5 mM MgCl₂, 20 pmol of primer, 500 pmol dNTP and 1 unit of Taq DNA polymerase (Perkin-Elmer, Branchburg, NJ). Forty cycles of amplification were performed using 94 °C denaturation for 30 seconds, 42 °C annealing for 1 minute and 72 °C extension for 30 seconds. Bands that were repeatedly observed to be specific to the RNA fingerprint pattern of the tumor were cut out of a silver stained gel, subcloned into the pGEM-T vector (Promega, Madison, WI) and sequenced. The isolated 3' sequences are provided in SEQ ID NO: 1-16.

> Comparison of these sequences to those in the public databases using the BLASTN program, revealed no significant homologies to the sequences provided in SEQ ID NO: 1-11. To the best of the inventors' knowledge, none of the isolated DNA sequences have previously been shown to be expressed at a greater level in human lung tumor tissue than in normal lung tissue.

aminopeptidase. Clone LT86-9 appears to contain two inserts, with the 5' sequence showing homology to the previously identified antisense sequence of interferon alpha-induced P27, and the 3' sequence being similar to LT86-6. Clone LT86-14 (SEQ ID NO: 30) was found to show some homology to the trithorax gene and has an "RGD" cell attachment sequence and a beta-Lactamase A site which functions in hydrolysis of penicillin. Clones LT86-1, LT86-2, LT86-4, LT86-5 and LT86-10 (SEQ ID NOS: 17, 18, 20, 21 and 26, respectively) were found to show homology to previously identified genes. A subsequently determined extended cDNA sequence for LT86-4 is provided in SEQ ID NO: 66, with the corresponding predicted amino acid sequence being provided in SEQ ID NO: 67.

Subsequent studies led to the isolation of five additional clones, referred to as LT86-20, LT86-21, LT86-22, LT86-26 and LT86-27. The determined 5' cDNA sequences for LT86-20, LT86-22, LT86-26 and LT86-27 are provided in SEQ ID NO: 68 and 70-72, respectively, with the determined 3' cDNA sequences for LT86-21 being provided in SEQ ID NO: 69. The corresponding predicted amino acid sequences for LT86-20, LT86-21, LT86-15 22, LT86-26 and LT86-27 are provided in SEQ ID NO: 73-77, respectively. LT86-22 and LT86-27 were found to be highly similar to each other. Comparison of these sequences to those in the gene bank as described above, revealed no significant homologies to LT86-22 and LT86-27. LT86-20, LT86-21 and LT86-26 were found to show homology to previously identified genes.

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predicted amino acid sequences are provided in SEQ ID NO: 93-101, respectively. L86S-30, L86S-39 and L86S-47 were found to be similar to each other. Comparison of these sequences with those in the gene bank as described above, revealed no significant homologies to L86S-14. L86S-29 was found to show some homology to a previously identified EST. L86S-6, L86S-11, L86S-34, L86S-39, L86S-47, L86S-49 and L86S-51 were found to show some homology to previously identified genes.

In further studies, a directional cDNA library was constructed using a Stratagene kit with a Lambda Zap Express vector. Total RNA for the library was isolated from two primary squamous lung tumors and poly A+ RNA was isolated using an oligo dT column. Antiserum was developed in normal mice using a pool of sera from three SCID mice implanted with human squamous lung carcinomas. Approximately 700,000 PFUs were screened from the unamplified library with *E. coli* absorbed mouse anti-SCID tumor serum. Positive plaques were identified as described above. Phage was purified and phagemid excised for 180 clones with inserts in a pBK-CMV vector for expression in prokaryotic or eukaryotic cells.

The determined cDNA sequences for 23 of the isolated clones are provided in SEQ ID NO: 126-148. Comparison of these sequences with those in the public database as described above revealed no significant homologies to the sequences of SEQ ID NO: 139 and 143-148. The sequences of SEQ ID NO: 126-138 and 140-142 were found to show homology previously identified human polynucleotide sequences.

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tags (ESTs). The sequences of SEQ ID NO: 150, 155 and 159-181 were found to show homology to sequences previously identified in humans.

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Example 6

ISOLATION OF DNA SEQUENCES ENCODING LUNG TUMOR ANTIGENS

DNA sequences encoding antigens potentially involved in squamous cell lung

5 tumor formation were isolated as follows:

A lung tumor directional cDNA expression library was constructed employing the Lambda ZAP Express expression system (Stratagene, La Jolla, CA). Total RNA for the library was taken from a pool of two human squamous epithelial lung carcinomas and poly A+ RNA was isolated using oligo-dT cellulose (Gibco BRL, Gaithersburg, MD). Phagemid were rescued at random and the cDNA sequences of isolated clones were determined.

The determined cDNA sequence for the clone SLT-T1 is provided in SEQ ID NO: 102, with the determined 5' cDNA sequences for the clones SLT-T2, SLT-T3, SLT-T5, SLT-T7, SLT-T9, SLT-T10, SLT-T11 and SLT-T12 being provided in SEQ ID NO: 103-110, respectively. The corresponding predicted amino acid sequence for SLT-T1, SLT-T2, SLT-T3, SLT-T10 and SLT-T12 are provided in SEQ ID NO: 111-115, respectively. Comparison of the sequences for SLT-T2, SLT-T3, SLT-T5, SLT-T7, SLT-T9 and SLT-T11 with those in the public databases as described above, revealed no significant homologies. The sequences for SLT-T10 and SLT-T12 were found to show some homology to sequences previously identified in humans.

The sequence of SLT-T1 was determined to show some homology to a PAC clone of unknown protein function. The cDNA sequence of SLT-T1 (SEQ ID NO: 102) was found to contain a mutator (MUTT) domain. Such domains are known to function in removal of damaged guanine from DNA that can cause A to G transversions (see, for example, el-Deiry, W.S., 1997 Curr. Opin. Oncol. 9:79-87; Okamoto, K. et al. 1996 Int. J. Cancer 65:437-41; Wu, C. et al. 1995 Biochem. Biophys. Res. Commun. 214:1239-45; Porter, D.W. et al. 1996 Chem. Res. Toxicol. 9:1375-81). SLT-T1 may thus be of use in the treatment, by gene therapy, of lung cancers caused by, or associated with, a disruption in DNA repair.

Example 7 SYNTHESIS OF POLYPEPTIDES

Polypeptides may be synthesized on a Perkin Elmer/Applied Biosystems Division 430A peptide synthesizer using FMOC chemistry with HPTU (O-Benzotriazole-N,N,N',N'-tetramethyluronium hexafluorophosphate) activation. A Gly-Cys-Gly sequence may be attached to the amino terminus of the peptide to provide a method of conjugation, binding to an immobilized surface, or labeling of the peptide. Cleavage of the peptides from the solid support may be carried out using the following cleavage mixture: trifluoroacetic acid:ethanedithiol:thioanisole:water:phenol (40:1:2:2:3). After cleaving for 2 hours, the peptides may be precipitated in cold methyl-t-butyl-ether. The peptide pellets may then be dissolved in water containing 0.1% trifluoroacetic acid (TFA) and lyophilized prior to purification by C18 reverse phase HPLC. A gradient of 0%-60% acetonitrile (containing 0.1% TFA) in water (containing 0.1% TFA) may be used to elute the peptides. Following lyophilization of the pure fractions, the peptides may be characterized using electrospray or other types of mass spectrometry and by amino acid analysis.

From the foregoing, it will be appreciated that, although specific embodiments of the invention have been described herein for the purposes of illustration, various modifications may be made without deviating from the spirit and scope of the invention.

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- 9. A vaccine comprising the polypeptide of claim 2 and an immune response enhancer.
- The vaccine of claim 9 wherein the immune response enhancer is an adjuvant.
 - 11. A vaccine comprising the polynucleotide of claims 1 or 4 and an immune response enhancer.

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- 12. The vaccine of claim 11 wherein the immune response enhancer is an adjuvant.
- 13. A pharmaceutical composition for the treatment of lung cancer comprising a polypeptide and a physiologically acceptable carrier, the polypeptide comprising an immunogenic portion of a lung protein or of a variant thereof, wherein said protein comprises an amino acid sequence encoded by a polynucleotide comprising a sequence selected from the group consisting of:
 - (a) sequences recited in SEQ ID NO: 12-18, 20, 21, 26, 49, 50, 52, 54, 64, 66, 68, 69, 71, 78, 84, 85, 88, 91, 92, 116-120, 126-138, 140-142, 150, 155 and 159-181;
 - (b) sequences complementary to the sequences of SEQ ID NO: 12-18, 20, 21, 26, 49, 50, 52, 54, 64, 66, 68, 69, 71, 78, 84, 85, 88, 91, 92, 116-120, 126-138, 140-142, 150, 155 and 159-181; and
 - (c) variants of the sequences of (a) and (b).

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- 14. A vaccine for the treatment of lung cancer comprising a polypeptide and an immune response enhancer, said polypeptide comprising an immunogenic portion of a lung protein or of a variant thereof, wherein said protein comprises an amino acid sequence encoded by a polynucleotide comprising a sequence selected from the group consisting of:
- (a) sequences recited in SEQ ID NO: 12-18, 20, 21, 26, 49, 50, 52, 54, 64, 66, 68, 69, 71, 78, 84, 85, 88, 91, 92, 116-120, 126-138, 140-142, 150, 155 and 159-181;

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- 21. A pharmaceutical composition comprising a fusion protein according to any one of claims 18-20 and a physiologically acceptable carrier.
- 5 22. A vaccine comprising a fusion protein according to any one of claims 18-20 and an immune response enhancer.
 - 23. The vaccine of claim 22 wherein the immune response enhancer is an adjuvant.

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- 24. A method for inhibiting the development of lung cancer in a patient, comprising administering to the patient an effective amount of the pharmaceutical composition of claim 21.
- 25. A method for inhibiting the development of lung cancer in a patient, comprising administering to the patient an effective amount of the vaccine of claim 22.
 - 26. A method for inhibiting the development of lung cancer in a patient, comprising administering to the patient a polynucleotide under conditions such that the polynucleotide enters a cell of the patient and is expressed therein, the polynucleotide having a sequence selected from the group consisting of:
 - (a) a sequence provided in SEQ ID NO: 102;
 - (b) sequences complementary to a sequence of SEQ ID NO: 102; and

- (c) variants of the sequence of SEO ID NO: 102.
- 27. A method for detecting lung cancer in a patient, comprising:
 - (a) contacting a biological sample obtained from the patient with a binding agent which is capable of binding to a polypeptide, the polypeptide comprising an immunogenic portion of a lung tumor protein or a variant thereof, wherein said protein comprises an amino acid sequence encoded by a polynucleotide comprising a nucleotide sequence selected from the group consisting of sequences provided in SEQ ID NO: 1-31, 49-

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- (a) sequences recited in SEQ ID NO: 1-11, 19, 22-25, 27-31, 51, 53, 55, 63, 70, 72, 79, 80, 86, 87, 89, 90, 102-107, 109, 139, 143-149, 151-154 and 156-158;
- (b) the complements of nucleotide sequences recited in SEQ ID NO: 1-11, 19, 22-25, 27-31, 51, 53, 55, 63, 70, 72, 79, 80, 86, 87, 89, 90, 102-107, 109, 139, 143-149, 151-154 and 156-158; and
- (c) variants of the sequences of (a) and (b).
- 32. A method for inhibiting the development of lung cancer in a patient, comprising administering to the patient a therapeutically effective amount of a monoclonal antibody according to claim 31.
- 33. The method of claim 32 wherein the monoclonal antibody is conjugated to a therapeutic agent.
 - 34. A method for detecting lung cancer in a patient comprising:
 - (a) obtaining a biological sample from the patient;
- (b) contacting the sample with at least two oligonucleotide primers in a polymerase chain reaction, wherein at least one of the oligonucleotides is specific for a polynucleotide encoding a polypeptide comprising an immunogenic portion of a lung tumor protein or a variant thereof, said protein comprising an amino acid sequence encoded by a polynucleotide comprising a nucleotide sequence selected from the group consisting of sequences recited in SEQ ID NO: 1-31, 49-55, 63, 64, 66, 68-72, 78-80, 84-92, 102-110, 116-120 and 126-181, the complements of said sequences and variants thereof; and
- (c) detecting in the sample a DNA sequence that amplifies in the presence of the oligonucleotide primers, thereby detecting lung cancer.
- 35. The method of claim 34, wherein at least one of the oligonucleotide primers comprises at least about 10 contiguous nucleotides of a polynucleotide comprising a sequence selected from SEQ ID NO: 1-31, 49-55, 63, 64, 66, 68-72, 78-80, 84-92, 102-110, 116-120 and 126-181.

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provided in SEQ ID NO: 1-31, 49-55, 63, 64, 66, 68-72, 78-80, 84-92, 102-110, 116-120 and 126-181, the complements of said sequences and variants thereof.

- 44. A method for detecting lung cancer in a patient, comprising:
- (a) obtaining a biological sample from the patient;
- (b) contacting the biological sample with an oligonucleotide probe specific for a polynucleotide encoding a polypeptide comprising an immunogenic portion of a lung tumor protein or a variant thereof, said protein comprising an amino acid sequence encoded by a polynucleotide comprising a nucleotide sequence selected from the group consisting of sequences recited in SEQ ID NO: 1-31, 49-55, 63, 64, 66, 68-72, 78-80, 84-92, 102-110, 116-120 and 126-181, the complements of said nucleotide sequences and variants thereof; and
- (c) detecting in the sample a DNA sequence that hybridizes to the oligonucleotide probe, thereby detecting lung cancer in the patient.
- 45. The method of claim 44 wherein the oligonucleotide probe comprises at least about 15 contiguous nucleotides of a polynucleotide having a nucleotide sequence selected from the group consisting of sequences recited in SEQ ID NO: 1-31, 49-55, 63, 64, 66, 68-72, 78-80, 84-92, 102-110, 116-120 and 126-181, the complements of said nucleotide sequences and variants thereof.
- A diagnostic kit comprising an oligonucleotide probe specific for a polynucleotide encoding a polypeptide comprising an immunogenic portion of a lung tumor protein or a variant thereof, said protein comprising an amino acid sequence encoded by a polynucleotide comprising a nucleotide sequence selected from the group consisting of sequences recited in SEQ ID NO: 1-31, 49-55, 63, 64, 66, 68-72, 78-80, 84-92, 102-110, 116-120 and 126-181, the complements of said sequences and variants thereof.
- 47. The diagnostic kit of claim 46, wherein the oligonucleotide probe comprises at least about 15 contiguous nucleotides of a polynucleotide having a nucleotide sequence selected from the group consisting of sequences recited in SEQ ID NO: 1-31, 49-55,

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pharmaceutically acceptable carrier.

- 55. A composition for the treatment of lung cancer in a patient, comprising T cells proliferated in the presence of a polynucleotide of claim 1, in combination with a pharmaceutically acceptable carrier.
 - 56. A method for treating lung cancer in a patient, comprising the steps of:
 - (a) incubating antigen presenting cells in the presence of at least one polypeptide of claim 2; and
 - (b) administering to the patient the incubated antigen presenting cells.
 - 57. A method for treating lung cancer in a patient, comprising the steps of:
 - (a) incubating antigen presenting cells in the presence of at least one polynucleotide of claim 1; and
 - (b) administering to the patient the incubated antigen presenting cells.
 - 58. The method of claims 54 or 55 wherein the antigen presenting cells are selected from the group consisting of dendritic cells and macrophage cells.
- 59. A composition for the treatment of lung cancer in a patient, comprising antigen presenting cells incubated in the presence of a polypeptide of claim 2, in combination with a pharmaceutically acceptable carrier.
- 60. A composition for the treatment of lung cancer in a patient, comprising antigen presenting cells incubated in the presence of a polynucleotide of claim 1, in combination with a pharmaceutically acceptable carrier.

SEQUENCE LISTING

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Leu	Asp	Ala	Gln	Glu	His	·Val	Lys	: Asn	Pro	Tyr	Lys	Gly	Lys	Lys	Leu	. Atra
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Tyr	Glu	Lys	Ala 180	Tyr	Glu	Gly	Gly	Ser 185	Glu	Leu	Gly		190			Ş4 177 - 174
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Gly	Arg	195	Thr	His	Asp	GIA	11e 200					205	Pne	PIO	Asp	Α.,
			7	T	T	C3	Ala	<u></u>					λla	Pro	Ara	
rea	210	Ser	Leu	Leu	11p	215	AIG	Gru	Азр	GIM	220	ASII	AIG	110	мg	·
Lys 225	Val	Pro	Asn	His	Tyr 230	Ile	Ala	Ile	Pro	Glu 235	Trp	Phe	Leu	Ser	Glu- 240	- (1) - (3) (1) - (3) (3)
). Den	'Al'a	Thr	Val	Δla	Thr	Glu	Thr	Ara	Ala	Val	Ile	λla	Tro	Met	Glu	ş
Moli	ΑIG	1111	Vai	245	*****	Ozu			250					255	-1.5	er i
īvs	Ile	Pro	Phe	Val	Leu	Gly	Gly	Asn	Leu	Gln	Gly	Gly	Glu	Leu	Val	15. B
			260			4		265			-		270			ŧ
Val	Ala	Тут 275	Pro	Tyr	Asp	Met	Val 280	Arg	Ser	Leu	Trp	Lys 285		Gln	Glu	
His	Thr 290	Pro	Thr	Pro	Asp	Asp 295	His	Val	Phe	Arg	Trp 300	Leu	Ala	Tyr	Ser	eeft z
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Tyr 305	Ala	Ser	Thr	.His	Arg 310	Leu	Met	Thr	Asp	Ala 315		Arg	Arg	Val	Cys 320	:
His	Thr	Glu	Asp	Phe-	Gln	Lys	Glu	Glu	Gly 330	Thr	Val	Asn	Gly	Ala 335	Ser	

Trp His Thr Val Ala Gly Ser Leu Asn Asp Phe Ser Tyr Leu His Thr 350. Asn Cys Phe Glu Leu Ser Ile Tyr Val Gly Cys Asp Lys Tyr Pro His 355 360 201 Glu Ser Glu Leu Pro Glu Glu Trp Glu Asn Asn Arg Glu Ser Leu Ile 370 Val Phe Met Glu Gln Val His Arg Gly Ile Lys Gly Ile Val Arg Asp 395 C. Leu Gln Gly Lys Gly Ile Ser Asn Ala Val Ile Ser Val Glu Gly Val Asn His Asp Ile Arg Thr Ala Ser Asp Gly Asp Tyr Trp Arg Leu Leu 430 Asn Pro Gly Glu Tyr Val Val Thr Ala Lys Ala Glu Gly Phe Ile Thr Ser Thr Lys Asn Cys Met Val Gly Tyr Asp Met Gly Ala Thr Arg Cys 450 grant 25 miles 455 days at the control 460 ft. Asp Phe Thr Leu Thr Lys Thr Asn Leu Ala Arg Ile Arg Glu Ile Met 465 480% Glu Thr Phe Gly Lys Gln Pro Val Ser Leu Pro Ser Arg Arg Leu Lys MEST CONTROL 485 ALM STREET AND CONTROL AND A 12 490 FOR CONTROL AND A 12 495 Leu Arg Gly Arg Lys Arg Arg Gln Arg Gly 18-10 500 - 18-2 12-2 18-2 18-505 - 1 <210> 35 · 25人的对方是 <211> 96 <212> PRT <400> 35 Met Asn Gly Glu Ala Asp Cys Pro Thr Asp Leu Glu Met Ala Ala Pro Arg Gly Gln Asp Arg Trp Ser Gln Glu Asp Met Leu Thr Leu Leu Glu 25 Cys Met Lys Asn Asn Leu Pro Ser Asn Asp Ser Ser Gln Phe Lys Thr 🦪 40 Thr Gln Thr His Met Asp Arg Glu Lys Val Ala Leu Lys Asp Phe Ser 55 50 Gly Asp Met Cys Lys Leu Lys Trp Val Glu Ile Ser Asn Glu Val Arg

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Lys Phe Arg Thr Leu Thr Glu Leu Ile Leu Asp Thr Gln Glu His Val 85 90 95

<210> 36

<211> 129

<212> PRT

<213> Homo sapiens

<400> 36

Gly Ile Val Val Phe Ser Leu Gly Ser Met Val Ser Glu Ile Pro Glu

1 5 10 15

Lys Lys Ala Val Ala Ile Ala Asp Ala Leu Gly Lys Ile Pro Gln Thr 20 25 30

Val Leu Trp Arg Tyr Thr Gly Thr Arg Pro Ser Asn Leu Ala Asn Asn 35 40 45

Thr Ile Leu Val Gln Trp Leu Pro Gln Asn Asp Leu Leu Gly His Pro 50 55 60

Met Thr Arg Ala Phe Ile Thr His Ala Ser Ser His Gly Val Asn Glu 65 70 75 80

Ser Ile Cys Asn Gly Val Pro Met Val Met Ile Pro Leu Phe Gly Asp 85 90 95

Gln Met Asp Asn Ala Lys Arg Arg Glu Thr Lys Gly Ala Gly Val Thr 100 105 110

Leu Asn Val Leu Glu Met Thr Ser Glu Asp Leu Glu Asp Ala Leu Lys 115 120 125

Ser

<210> 37

<211> 238

<212> PRT

<213> Homo sapiens

<400> 37

Asn Leu Leu Gly Ile Ser Trp Val Asp Ser Ser Trp Ile Pro Ile Leu

1 5 10 15

Asn Ser Gly Ser Val Leu Asp Tyr Phe Ser Glu Arg Ser Asn Pro Phe 20 25 30

Tyr Asp Arg Thr Cys Asn Asn Glu Val Val Lys Met Gln Arg Leu Thr
35 40 45

Leu Glu His Leu Asn Gln Met Val Gly Ile Glu Tyr Ile Leu Leu His 50 55 60

Ala Gln Glu Pro Ile Leu Phe Ile Ile Arg Lys Gln Gln Arg Gln Ser 75 70 Pro Ala Gln Val Ile Pro Leu Ala Asp Tyr Tyr Ile Ile Ala Gly Val 90 Ile Tyr Gln Ala Pro Asp Leu Gly Ser Val Ile Asn Ser Arg Val Leu 105 110 Thr Ala Val His Gly Ile Gln Ser Ala Phe Asp Glu Ala Met Ser Tyr Cys Arg Tyr His Pro Ser Lys Gly Tyr Trp Trp His Phe Lys Asp His 135 130 Glu Glu Gln Asp Lys Val Arg Pro Lys Ala Lys Arg Lys Glu Glu Pro Ser Ser Ile Phe Gln Arg Gln Arg Val Asp Ala Leu Leu Leu Asp Leu 170 165 Arg Gln Lys Phe Pro Pro Lys Phe Val Gln Leu Lys Pro Gly Glu Lys 185 190 Pro Val Pro Val Asp Gln Thr Lys Lys Glu Ala Glu Pro Ile Pro Glu 200 205 195 Thr Val Lys Pro Glu Glu Lys Glu Thr Thr Lys Asn Val Gln Gln Thr 215 220 Val Ser Ala Lys Gly Pro Pro Glu Lys Arg Met Arg Leu Gln 230 225 <210> 38 <211> 202 <212> PRT <213> Homo sapiens <400> 38 Lys Gly Ser Glu Gly Glu Asn Pro Leu Thr Val Pro Gly Arg Glu Lys 1 Glu Gly Met Leu Met Gly Val Lys Pro Gly Glu Asp Ala Ser Gly Pro Ala Glu Asp Leu Val Arg Arg Ser Glu Lys Asp Thr Ala Ala Val Val 40 Ser Arg Gln Gly Ser Ser Leu Asn Leu Phe Glu Asp Val Gln Ile Thr 55 Glu Pro Glu Ala Glu Pro Glu Ser Lys Ser Glu Pro Arg Pro Pro Ile 75

Ser Ser Pro Arg Ala Pro Gln Thr Arg Ala Val Lys Pro Arg Leu His 85 90 95

Pro Val Lys Pro Met Asn Ala Thr Ala Thr Lys Val Ala Asn Cys Ser 100 105 110

Leu Gly Thr Ala Thr Ile Ile Gly Glu Asn Leu Asn Asn Glu Val Met
115 120 125

Met Lys Lys Tyr Ser Pro Ser Asp Pro Ala Phe Ala Tyr Ala Gln Leu 130 135 140

Thr His Asp Glu Leu Ile Gln Leu Val Leu Lys Gln Lys Glu Thr Ile 145 150 155 160

Ser Lys Lys Glu Phe Gln Val Arg Glu Leu Glu Asp Tyr Ile Asp Asn 165 170 175

Leu Leu Val Arg Val Met Glu Glu Thr Pro Asn Ile Leu Arg Ile Pro 180 185

Thr Gln Val Gly Lys Lys Ala Gly Lys Met
195 200

<210> 39

<211> 243

<212> PRT

<213> Homo sapiens

<400> 39

Val Asn Ala Leu Gly Ile Met Ala Ala Val Asp Ile Arg Asp Asn Leu

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Leu Gly Ile Ser Trp Val Asp Ser Ser Trp Ile Pro Ile Leu Asn Ser 20 25 30

Gly Ser Val Leu Asp Tyr Phe Ser Glu Arg Ser Asn Pro Phe Tyr Asp
35
40
45

Arg Thr Cys Asn Asn Glu Val Val Lys Met Gln Arg Leu Thr Leu Glu
50 55 60

His Leu Asn Gln Met Val Gly Ile Glu Tyr Ile Leu Leu His Ala Gln 65 70 75 80

Glu Pro Ile Leu Phe Ile Ile Arg Lys Gln Gln Arg Gln Ser Pro Ala 85 90 95

Gln Val Ile Pro Leu Ala Asp Tyr Tyr Ile Ile Ala Gly Val Ile Tyr 100 105 110

Gln Ala Pro Asp Leu Gly Ser Val Ile Asn Ser Arg Val Leu Thr Ala 115 120 125

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۷a	1 Hi 13	s Gl O	y Il	e Gl	n Se	r Ala 13		e As	p Gl	u Ala	a Met 140		Туг	Cys	s Arg	
Ту: 14!		s Pr	o Se	r Ly	s Gly 15		r Tr	o Tr	p Hi	s Phe 159		Ası) His	Glu	1 Glu 160	e, See e
Glı	n As	р Lу	s Va	1 Arg		o Lys	s Ala	а Гу	s Ar		g Glu	Glu	Pro	Ser 175	Ser	e E Ng
Ile	e Ph	e Gla	n Arg		n Arg	g V al	Asp	18		u Leu	Leu	Asp	Leu 190		(Gln	i suiji
Lys	s Ile	e Se:	r Thi	c Glr	n Ile	e Cys	Ala 200		l Asp	Gln	Thr	Lys 205	-	Glu	Ala	3-76 8€:.
Glu	210	o Ile	e Pro	Glu	ı Thr	Val	Lys	Pro	o Gl u	ı Gİu	Lys 220		Thr	Thr	Lys	
Asr 225	ı_Va	l Gli	ı Glr	1 Thr	230		Ala	Lys	s Gly	Pro 235		G1u	Lys	Arg	Met 240	t żwit
Arg	Let	ı Glr	1			•	n. i	is‴s⊤, •		 .*	٠.	. ·•	: 24		19.2.27 19	
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			Asp	Ile 5		Asp	Asn	Leu	Leu 10		Ile	Ser	Trp	Val		្រាស់ ស្រីស ប្រាស់ ស្រីស់ ទីស
er	Ser	Trp	Ile 20		Ile	Leu	Asn	Ser 25	Gly		Val	Leu	Asp 30	Tyr		
er	Glu	Arg 35		Asn	Pro	Phe	Tyr 40		Arg	Thr	Cys	Asn 45	Asn	Glu	Val	
al	Lys 50	Met	Gln	Arg	Leu	Thr 55	Leu	Glu	His	Leu	Asn 60	Gln	Met	Val	Gly	इ. हा क
1e 65	Glu	Tyr	Ile	Leu	Leu 70	His	Ala	Gln	Glu						Ile 80	
rg	Lys	Gln	Gln	Arg 85	Gln	Ser	Pro		Gln 90						Asp	ad Turk
yr	Tyr	Ile	Ile 100		Gly	Val	Ile	Tyr		Ala					Ser	5 / CF
al	Ile	Asn 115		Arg	Val	Leu	Thr 120		Val	His		Ile	Gln	Ser	Ala	

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Phe	Asp 130	Glu	Ala	Met	Ser	135	Cys	Arg	ЛУĽ	HIS	140	ser	rys	GIÀ	lyr
Trp	Trp	His	Phe	Lys	Asp 150	His	Glu	Glu		Asp 155	Lys	V al	Arg	Pro	Lys 160
Ala	Lys	Arg	Lys	Glu 165	Glu	Pro	Ser	Ser	Ile 170	Phe	Gln	Arg	Gln	Arg 175	Val
Asp	Ala	Leu	Leu 180	Leu	Asp	Leu	Arg	Gln 185	Lys	Phe			Lys 190	7.75	Val
Gln	Leu	Lys 195		Gly	Glu	Lys	Pro 200	Val	Pro		Asp	205		Lys	Lys
Glu	Ala 210		Pro	Ile	Pro	Glu 215	Thr	Val	Lys					Glu	Thr
Thr 225	Lys	Asn	'Val'	Gln	Gln 230	Thr	Val.	Ser		235	Gly	Pro	Pro	Glu	Lys 240
Arg	Met	Arg		G1n 245	*				· .	•		•		ega j	-
<211)> 41 l> 16 2> PF	53			ista. N		<i>f</i>		April 1	. '` ``	file o	. 1 . 4)		an T	: . 7. ·
<213			ء هــــــــــــــــــــــــــــــــــــ		12 2	se s .							:		
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130

135

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Val 145		Leu	Pro	Arg	Pro 150		Gln	Asp	Leu	Leu 155		His	Glu	. Ser	Let 160
Leu	Ala	Ala								. •				n e	
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	2> P						1.					1			•
<21	3> H	omo	sapi	ens	j.fr.					j		. 37.	. Sa . Sa .		
<40	0> 4	્ં 2		1 (187) 		•		. :.							•
			Arg	Asp	Asn	Leu	Leu	Gly	Ile	Ser	Trp	Val	Asp	Ser	Ser
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					Asn	Ser	Gly	Ser 25	Val	Leu	Asp	Tyr		Ser	Glu
	왕. 발.		. 20					23	٠				30		•
		Asn	Pro	Phe	Tyr	Asp	Arg	Thr	Cys	Asn	Asn	Glu	Val	Val	Lys
		35	•				40		-			45			
Met	Gln 50	Arg	Leu	Thr	Leu	Glu 55	His	Leu	Asn	Gln	Met 60	Val	Gly	Ile	Glu
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Tyr 65	Ile	Leu	Leu	His	70	Gln	Glu	Pro	Ile	Leu 75	Phe	Ile	Ile	Arg	Lys 80
Gln	Gln	Arg		Ser 85	Pro	Ala	Gln	Val	Ile 90	Pŗo	Leu	Ala	Asp	Tyr 95	Туг
Ile	Ile		Gly 100	Val	Ile	Tyr	Gln	Ala 105	Pro	Asp	Leu	* *** ***	Ser 110	.Va1	Ile
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Asn	Ser				Thr				Gly	Ile	Gln	Ser 125	Ala	Phe	Asp
Glu	Ala	Met	Ser	Tyr	Cys	Arg	Tyr	Hiş	Pro	Ser	Lys	Gly	Tyr	Trp	Trp
	130	. 1.	1.3	Ž.	Service Control	135		.**.	•		140				,
	Phe	Lys	Asp	His		Glu	Gln	Asp	Lys	145 51 51	Arg	Pro	Lys	Ala	
145	15		,		150	<i>:</i> .				155	. •				1,60
Arg	Lys	Glu	Glu	Pro	Ser	Ser	Ile	Phe	Gln	Arg	Gln	Arg	Val	Asp	Ala
		. 1 21		165	1.				170			42	-	175	
Leu		Leu		Leu	• .	Gln	Lys	Phe 185	Pro	Pro	Lys	Phe	Val 190		Leu
_						_									
Lys	Pro	Gly 195	Glu	Lys	Pro		Pro 200	Val	Asp	Gln	5 - 1	Lys 205	Lys	Glu	Ala
Glu	Pro 210	Ile		Glu		Val 215	Lys	Pro	Glu	Glu	Lys 220	Glu	Thr	Thr	Lys

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Arg Leu Gln

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Ser Trp Ile Pro Ile Leu Asn Ser Gly Ser Val Leu Asp Tyr Phe Ser

Glu Arg Ser Asn Pro Phe Tyr Asp Arg Thr Cys Asn Asn Glu Val Val 35 40 45

∂ 25.__

Lys Met Gln Arg Leu Thr Leu Glu His Leu Asn Gln Met Val Gly Ile
50 60

Glu Tyr Ile Leu Leu His Ala Gln Glu Pro Ile Leu Phe Ile Ile Arg
65 70 75 7 80

Lys Gln Gln Arg Gln Ser Pro Ala Gln Val Ile Pro Leu Ala Asp Tyr

Tyr Ile Ile Ala Gly Val Ile Tyr Gln Ala Pro Asp Leu Gly Ser Val

Ile Asn Ser Arg Val Leu Thr Ala Val His Gly Ile Gln Ser Ala Phe 115 120 125

Asp Glu Ala Met Ser Tyr Cys Arg Tyr His Pro Ser Lys Gly Tyr Trp

Trp His Phe Lys Asp His Glu Glu Gln Asp Lys Val Arg Pro Lys Ala (1986) 145

Lys Arg Lys Glu Glu Pro Ser Ser Ile Phe Gln Arg Gln Arg Val Asp

Leu Lys Pro Gly Glu Lys Pro Val Pro Val Asp Gln Thr Lys Lys Glu 195

Lys Asn Val Gln Gln Thr Val Ser Ala Lys Gly Pro Pro Glu Lys Arg 225 · 230 235 Met Arg Leu Gln <210> 44 <211> 109 <212> PRT <213> Homo sapiens <400> 44 Glu Leu His Phe Ser Glu Phe Thr Ser Ala Val Ala Asp Met Lys Asn 1 Ser Val Ala Asp Arg Asp Asn Ser Pro Ser Ser Cys Ala Gly Leu Phe Ile Ala Ser His Ile Gly Phe Asp Trp Pro Gly Val Trp Val His Leu 40 Asp Ile Ala Ala Pro Val His Ala Gly Glu Arg Ala Thr Gly Phe Gly Val Ala Leu Leu Ala Leu Phe Gly Arg Ala Ser Glu Asp Pro Leu Leu Asn Leu Val Ser Pro Leu Asp Cys Glu Val Asp Ala Gln Glu Gly Asp Asn Met Gly Arg Asp Ser Lys Arg Arg Arg Leu Val 100 105

<210> 45 <211> 324 <212> PRT <213> Homo sapiens

ा के अनुस्कार रहते. एको उपने अनुस्कार है के अनुस्कार अनुस्कार करने हैं। Arg Arg Pro Val Met Ala Gln Glu Thr Ala Pro Pro Cys Gly Pro Val

Ser Arg Gly Asp Ser Pro Ile Ile Glu Lys Met Glu Lys Arg Thr Cys 25

Ala Leu Cys Pro Glu Gly His Glu Trp Ser Gln Ile Tyr Phe Ser Pro 40

Ser Gly Asn Ile Val Ala His Glu Asn Cys Leu Leu Tyr Ser Ser Gly 55

Leu Val Glu Cys Glu Thr Leu Asp Leu Arg Asn Thr Ile Arg Asn Phe

Asp Val Lys Ser Val Lys Lys Glu Ile Trp Arg Gly Arg Arg Leu Lys 85 90 95

Cys Ser Phe Cys Asn Lys Gly Gly Ala Thr Val Gly Cys Asp Leu Trp
100 105 110

Phe Cys Lys Lys Ser Tyr His Tyr Val Cys Ala Lys Lys Asp Gln Ala 115 120 125

Ile Leu Gln Val Asp Gly Asn His Gly Thr Tyr Lys Leu Phe Cys Pro 130 135 140

Glu His Ser Pro Glu Gln Glu Glu Ala Thr Glu Ser Ala Asp Asp Pro 145 150 155 160

Ser Met Lys Lys Lys Arg Gly Lys Asn Lys Arg Leu Ser Ser Gly Pro 165 170 175

Pro Ala Gln Pro Lys Thr Met Lys Cys Ser Asn Ala Lys Arg His Met
180 185 190

Thr Glu Glu Pro His Gly His Thr Asp Ala Ala Val Lys Ser Pro Phe
195 200 205

Leu Lys Lys Cys Gln Glu Ala Gly Leu Leu Thr Glu Leu Phe Glu His 210 215 220

1. 1. 1

Ile Leu Glu Asn Met Asp Ser Val His Gly Arg Leu Val Asp Glu Thr 225 230 235 240

Ala Ser Glu Ser Asp Tyr Glu Gly Ile Glu Thr Leu Leu Phe Asp Cys 245 250 255

Gly Leu Phe Lys Asp Thr Leu Arg Lys Phe Gln Glu Val Ile Lys Ser 260 265 270

Lys Ala Cys Glu Trp Glu Glu Arg Gln Arg Gln Met Lys Gln Gln Leu 275 280 285

Glu Ala Leu Ala Asp Leu Gln Gln Ser Leu Cys Ser Phe Gln Glu Asn 290 295 300

Gly Asp Leu Asp Cys Ser Ser Ser Thr Ser Gly Ser Leu Leu Pro Pro 305 310 315 320

Glu Asp His Gln

<210> 46

<211> 244

<212> PRT

<213> Homo sapiens

<400> 46

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Glu	Arg		Asn		Phe	Tyr	Asp 40	Arg	Thr	Cys	Asn	A sn 45	Glu	Val	Val
Lys	Met 50	Gln	Arg	Leu	Thr	Leu 55	Glu	His	Leu	Asn	Gln 60	Met	Val	Gly	Ile
Glu 65	Tyr	Ile	Leu	Leu	His 70	Ala	Gln	Glu				Phe			
Lys	Gln	Gln	Arg	Gln 85	Ser	Pro	Ala	Gln	Val 90	Ile	Pro	Leu	Ala	Asp 95	Tyr
_	Ile		100	··		4	41.4	105	5.4		i i i i i i i i i i i i i i i i i i i		110	f exit	
•		115				·	120			·	· 필요함	125		·	
	Glu 130		•	:	٠	135		*	· .	i kari Bud	140	1 41	Ç.	,	
145	*		- T. 144.		150		e.	• •		155	v Gran	, * · · · · · · · · · ·	,		160
	Arg		i"	165	. ,	. 21	. : 5		170		,	***,		175	•
	Leu	• • •	180			•		185					190		÷.
		195	,	·		* 1	200			· · ·	-	205	,		:
	Glu 210					215			s î.		220		- మెమ్	·•	
Lys 225	Asn	Val	Gln	Gln	Thr 230	Val	Ser	Ala (Lys	Gly 235	Pro	Pro	Glu	Lys	Arg 240
Met	Arg	Leu	Gln								,				

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   cgaacgcggc.tcgaatggca agccaaaatt ccttccggat agaatatgat acctttggtg: 180
  aactaaaggt gccaaatgat aagtattatg gcgcccagac cgtgagatct acgatgaact 240
  tgaagegage ggcegetgaa gtaaaceagg attatggtet tgateeaaag attgetaatg 360
  caataatgaa ggcagcagat gaggtagctg aaggtaaatt aaatgatcat tttcctctcg 420
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  aggaaataga agttgcccca ccaaagacta aagaagttcg cattaagatt ttggccacag 180
  gaatctgtcg cacagatgac catgtgataa aaggaacaat ggtgtccaag tttccagtga 240
  ttgtgggaca tgaggcaact gggattgtag agagcattgg agaaggagtg actacagtga 300
  aaccaggtga caaagtcatc cotototto tgccacaatg tagagaatgc aatgottgto 360
  gcaacccaga tggcaacctt tgcattagga gcgatattac tggtcgtgga gtactggctg 420
  atggcaccac cagatttaca tgcaagggcg aaccagtcca ccacttcatg aacaccagta 480
g catttacoga gtacacagt | respectively belong the symbol of the second respectively and a 499
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ada omi sare. T
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  <211> 887
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  aaagtggcag agetgtatte tatecataae tetggagaea aatetgatat teaggaeete 180
  ctggagagtg tcaggctgga caaagaaaaa gcagagactt tggctagtag cttgcaggaa 240
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 ttagaaaaat taagatcaga cctggatgaa aaagaaacag aaaggagtga catgaaagaa 420
 accatetttg aacttgaaga tgaagtagaa caacategtg etgtgaaact teatgacaac 480
  ctcattattt ctgatctaga gaatacagtt aaaaaactcc aggaccaaaa gcacgacatg 540
```

```
gaaagagaaa taaagacact ccacagaaga Cttcgggaag aatctgcgga atggcggcag 600
tttcaggctg atctccagac tgcagtagtc attgcaaatg acattaaatc tgaagcccaa 660
gaggagattg gtgatctaaa gcgccggtta catgaggctc aagaaaaaaa tgagaaactc 720
acaaaagaat tggaggaaat aaagtcacgc aagcaagagg aggagcgagg cgggtataca 780
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cctgctatag ctcagttttc agttcagaaa gtcactcctc agtctgatgg ctccagttca 180
aaagtgaaag tcaaagttog agtaaatgto catggcattt Lcagtgtgtc cagtgcatct 240
ttagtggagg ttcacaagtc tgaggaaaat gaggagccaa tggaaacaga tcagaatgca 300
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cagacaccag gcagaaaata aggcagagtc tgaagaaatg gagacctctc aagctggatc. 420
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tgtggacctg g
<210> 53
                                                            <211> 787
                                                           King a tuké
<212> DNA
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caggggtagt gatectggea gteaceatag efetaettgt ttaettttta gettftgate 180
aaaaatetta ettttatagg ageagtttte aacteetaaa tgttgaatat aatagteagt 240
taaattcacc agctacacag gaatacagga ctttgagtgg aagaattgaa tctctgatta 300
Ctaaaacatt caaagaatca aatttaagaa atcagttcat cagagctcat gttgccaaac 360
tgaggcaaga tggtagtggt gtgagagcgg atgttgtcat gaaatttcaa ttcactagaa 420
ataacaatgg agcatcaatg aaaagcagaa ttgagtctgt tttacgacaa atgctgaata 480
actetggaaa eetggaaata aaccetteaa etgagataac ateaettaet gaceaggetg 540
cagcaaattg gcttattaat gaatgtgggg ccggtccaga cctaataaca ttgtctgagc 600
agagaateet tggaggeaet gaggetgagg agggaagetg geegtggeaa gteagtetge 660
ggctcaataa tgcccaccac tgtggaggca gcctgatcaa taacatgtgg atcctgacag 720
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                                                          7.6%
ccacaac
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<213> Homo sapiens
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gaaccacatg ttgaagagca acagcagcag acaccagcag aaaataaggc agagtitgaa 180
gaaatggaga ceteteaage tggatecaag gataaaaaga tggaceaace acceeaagee 240
aagaaggcaa aagtgaagac cagtactgtg gacctgccaa tcgagaatca gctattatgg 300
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caggggtagt gatectggca gtcaccatag ctctacttgt ttacttttta gcttttgate 180
aaaaatctta cttttatagg agcagttttc aactcctaaa tgttgaatat aatagtcagt 240
taaattcacc agctacacag gaatacagga ctttgagtgg aagaattgaa tctctgatta 300
ctaaaacatt caaagaatca aatttaagaa atcagttcat cagagctcat gttgccaaac 360
tgaggcaaga tggtagtggt gtgagagcgg atgttgtcat gaaatttcaa ttcactagaa 420
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cageteactg etteagaage aactetaate etegtgactg gattgecaeg tetggtattt 780
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ctgttgcaaa gtctgtatgc aggtgtgcct gtcttaaatt ccaaagcttt acatttcaac 1380
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                                     10
Arg Ala Pro Ala Ala Ala Leu Ala Ser Ala Pro Gly Leu Gly Gly Ala
                                 25
Ala Val Pro Ser Phe Trp Pro Pro Asn Ala Ala Arg Met Ala Ser Gln
                                                45
       35
Asn Ser Phe Arg Ile Glu Tyr Asp Thr Phe Gly Glu Leu Lys Val Pro
Asn Asp Lys Tyr Tyr Gly Ala Gln Thr Val Arg Ser Thr Met Asn Phe
                                         75
 65
```

Lys Ile Gly Gly Val Thr Glu Arg Met Pro Thr Pro Val Ile Lys Ala 90

Phe Gly Ile Leu Lys Arg Ala Ala Ala Glu Val Asn Gln Asp Tyr Gly 105

Leu Asp Pro Lys Ile Ala Asn Ala Ile Met Lys Ala Ala Asp Glu Val 120

Ala Glu Gly Lys Leu Asn Asp His Phe Pro Leu Val Val Trp Gln Thr

Gly Ser Gly Thr Gln Thr Asn Met Asn Val Asn Glu Val Ile Ser 3.03 150

ကြောင်းကြောင့်များကြောင်းကြုံသည်။ ကြို့နောက်ကြောင်းမြောင်းများ

(2.10) graphs of the results defined by $\frac{1}{2}$. We have the regulation μ_{1}

<212> PRT <213> Homo sapiens

Lys Lys Ser Met Phe Ala Glu Ile Gln Ile Gln Asp Lys Asp Arg Met

A THE CHARLET DELLÉGIQUES É QUI MAÇIN A ÇA CATTERAT DE L'

Gly Thr Ala Gly Lys Val Ile Lys Cys Lys Ala Ala Val Leu Trp Glu

Gln Lys Gln Pro Phe Ser Ile Glu Glu Ile Glu Val Ala Pro Pro Lys

Thr Lys Glu Val Arg Ile Lys Ile Leu Ala Thr Gly Ile Cys Arg Thr

Asp Asp His Val Ile Lys Gly Thr Met Val Ser Lys Phe Pro Val Ile 75

Val Gly His Glu Ala Thr Gly Ile Val Glu Ser Ile Gly Glu Gly Val 95

Thr Thr Val Lys Pro Gly Asp Lys Val Ile Pro Leu Phe Leu Pro Gln

Cys Arg Glu Cys Asn Ala Cys Arg Asn Pro Asp Gly Asn Leu Cys Ile

Arg Ser Asp Ile Thr Gly Arg Gly Val Leu Ala Asp Gly Thr Thr Arg 135 140

Phe Thr Cys Lys Gly Glu Pro Val His His Phe Met Asn Thr Ser Thr 150 155

Phe Thr Glu Tyr Thr

165

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	11> 2				•							-			
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<21	13> · I	Omo	sapi	ens											
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Glu	ı Sei	Gli	ı Glr	Lvs	Glv	LVS	: Ala	Ala	Leu	Ala	Ala				
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Ala	Glr	Leu	ı Glu	Asn	Glu	Lys	Gln	Lys	Val	Ala	Glu	Leu	Tyr	Ser	Ile
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Hig	Agr		Gly												
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	50					55		_	-		60				
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		Asp	Lys	Glu	Lys	Ala	Glu	Thr	Leu	Ala	Ser	Ser	Leu	Gln	Glu
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Asp	Leu	Ala	His	Thr	Arg	Asn	Asp	Ala	Asn	Arq	Leu	Gln	Aso	Ala	Ile
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	٠.										والمراجع المراجع	·			
Δla	Laze	Val	Glu	Nen	Gly	Time	7~~	λla	Dhe	Cla	C1.	C1	33 (3.2) 3 3 3 3	Tara	T
	LJy 3	· •aı			Giu	IYL	AL 9			GII	GIU			гÀг	Lys
			100					105					.110		
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Gin	Ile		Asp	Leu	Asn	Met	Thr	Leu	Glu	Lys	Leu	Arg	Ser	Asp	Leu
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Asp	Glu	Lys	Glu												
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Leu	Ile	Ile	Ser	Asp	Leu	Glu	Asn	Thr	Val	Lys	Lys	Leu	Gln	Asp	Gln
				165					170					175	• :
	Sec .	. C		1 600	s 193.	9	a att	ř	5 .74 ⁷	2.353	V	: 55	i ov		
Lys	His	Asp	Met	Glu	Arg	Glu	Ile	Lys	Thr	Leu	His	Arg	Arg	Leu	Arq
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Glu	Gliv	Car	או ה	C1	T -	A	C3-	Dho	C1-	NI A	3	7	<i>~</i> 1~	/ /·	31-
GIU	Gru		Ala	GIU	пр	мy	GIII	Pile	GIII	Ald	ASD		GTII	THE	Ala
		195					200					205			
			1.1%												
Val	Val	Ile	Ala	Asn	Asp	Ile	Lys	Ser	Glu	Ala	Gln	Glu	Glu	Ile	Gly
	210					215			•		220				-
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Men			Arg	λ~~			Glie	λ1 -	Gla	G]	Terr	70 0	C1		T
	<u> </u>	nys	AL Y	мy		urz	GIU	vra	GIII		πλa	wall	GIU	пĀģ	
225					230					235					240
		. `:-	No grant							;**,	1.32	· 1,	5 1.5.4		: 7.
Thr	Lys	Glu	Leu	Glu	Glu	Ile	Lys	Ser	Arg	Lys	Gln	Glu	Glu	Glu	Arg
				245					250					255	-

Gly Gly Tyr

<210> 59

<211> 125

<212> PRT

<213> Homo sapiens

₹400> 59

Gly Thr Ser Phe Ser Lys Asn His Ala Ala Pro Phe Ser Lys Val Leu

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Thr Phe Tyr Arg Lys Glu Pro Phe Thr Leu Glu Ala Tyr Tyr Ser Ser 20 25 30

Pro Gln Asp Leu Pro Tyr Pro Asp Pro Ala Ile Ala Gln Phe Ser Val

Gln Lys Val Thr Pro Gln Ser Asp Gly Ser Ser Ser Lys Val Lys Val
50 60

Lys Val Arg Val Asn Val His Gly Ile Phe Ser Val Ser Ser Ala Ser 65

Leu Val Glu Val His Lys Ser Glu Glu Asn Glu Glu Pro Met Glu Thr

Asp Gln Asn Ala Lys Glu Glu Glu Lys Met Gln Val Asp Gln Glu Glu
100 105 110

Pro His Val Glu Glu Gln Gln Gln Gln Thr Pro Gly Arg
115 120 125

一声 化聚戊烯醇 医髓囊病 化基氯化 翻門 化二甲烷烷基

<210> 60

<211> 246@46 0... \ 1793 @\ 0...

<212> PRT

<213> Homo sapiens

<400> 60

Met Tyr Arg Pro Ala Arg Val Thr Ser Thr Ser Arg Phe Leu Asn Pro

Tyr Val Val Cys Phe Ile Val Val Ala Gly Val Val Ile Leu Ala Val
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Thr Ile Ala Leu Leu Val Tyr Phe Leu Ala Phe Asp Gln Lys Ser Tyr

Phe Tyr Arg Ser Ser Phe Gln Leu Leu Asn Val Glu Tyr Asn Ser Gln 50 55 60

Leu Asn Ser Pro Ala Thr Gln Glu Tyr Arg Thr Leu Ser Gly Arg Ile
65 70 75 80

Glu Ser Leu Ile Thr Lys Thr Phe Lys Glu Ser Asn Leu Arg Asn Gln
85 90 95

Phe Ile Arg Ala His Val Ala Lys Leu Arg Gln Asp Gly Ser Gly Val

Arg Ala Asp Val Val Met Lys Phe Gln Phe Thr Arg Asn Asn Gly
115 120 125

Ala Ser Met Lys Ser Arg Ile Glu Ser Val Leu Arg Gln Met Leu Asn 130 135 140

Asn Ser Gly Asn Leu Glu Ile Asn Pro Ser Thr Glu Ile Thr Ser Leu 145 150 155 160

Pro Asp Leu Ile Thr Leu Ser Glu Gln Arg Ile Leu Gly Gly Thr Glu
180 185 190

Ala Glu Glu Gly Ser Trp Pro Trp Gln Val Ser Leu Arg Leu Asn Asn 195 200 205

Ala His His Cys Gly Gly Ser Leu Ile Asn Asn Met Trp Ile Leu Thr 210 225

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Thr Ser Gly Ile Ser Thr

<210> 61

<211> 128

<212> PRT

<213> Homo sapiens

<400> 61

Gly Ile Phe Ser Val Ser Ser Ala Ser Leu Val Glu Val His Lys Ser 1 5 10 15

Glu Glu Asn Glu Glu Pro Met Glu Thr Asp Gln Asn Ala Lys Glu Glu

Glu Lys Met Gln Val Asp Gln Glu Glu Pro His Val Glu Gln Gln Gln 35 40 45

Gln Gln Thr Pro Ala Glu Asn Lys Ala Glu Ser Glu Glu Met Glu Thr

Ser Gln Ala Gly Ser Lys Asp Lys Lys Met Asp Gln Pro Pro Gln Ala 65 70 75 80

Lys Lys Ala Lys Val Lys Thr Ser Thr Val Asp Leu Pro Ile Glu Asn

. 85 90 95

Gln Leu Leu Trp Gln Ile Asp Arg Glu Met Leu Asn Leu Tyr Ile Glu 100 105 110

Asn Glu Gly Lys Met Ile Met Gln Asp Lys Leu Glu Lys Glu Arg Asn 115 120 125

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<211> 418

<212> PRT

<213> Homo sapiens

<400> 62

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Tyr Val Val Cys Phe Ile Val Val Ala Gly Val Val Ile Leu Ala Val

Thr Ile Ala Leu Leu Val Tyr Phe Leu Ala Phe Asp Gln Lys Ser Tyr. 35 40 45

Phe Tyr Arg Ser Ser Phe Gln Leu Leu Asn Val Glu Tyr Asn Ser Gln 50 55

Leu Asn Ser Pro Ala Thr Gln Glu Tyr Arg Thr Leu Ser Gly Arg Ile
65 70 75 80

Glu Ser Leu Ile Thr Lys Thr Phe Lys Glu Ser Asn Leu Arg Asn Gln
85 ... 90 95

Phe Ile Arg Ala His Val Ala Lys Leu Arg Gln Asp Gly Ser Gly Val 100 105 110

Arg Ala Asp Val Val Met Lys Phe Gln Phe Thr Arg Asn Asn Asn Gly
115 120 125

Ala Ser Met Lys Ser Arg Ile Glu Ser Val Leu Arg Gln Met Leu Asn 130 135 140

Asn Ser Gly Asn Leu Glu Ile Asn Pro Ser Thr Glu Ile Thr Ser Leu 145 150 155 160

Thr Asp Gln Ala Ala Asn Trp Leu Ile Asn Glu Cys Gly Ala Gly
165 170 175

Pro Asp Leu Ile Thr Leu Ser Glu Gln Arg Ile Leu Gly Gly Thr Glu 180 185 190

Ala Glu Glu Gly Ser Trp Pro Trp Gln Val Ser Leu Arg Leu Asn Asn 195 200 205

Ala His His Cys Gly Gly Ser Leu Ile Asn Asn Met Trp Ile Leu Thr

		210				1	215					220				-	
			4	• :,						14.			200	χ		• • • • • •	
	4	Ala	His	Cys	Phe	Arg	Ser	Asn	Ser	Asn	Pro	Arg	Asp	Trp	Ile	Ala	. *
	225			. *	: :	230	귝 .		7		235	•	. · · ·	e 100		240	-
				5.	<u>.</u>	٠,,								or is	. :	÷	•
•	Thr	Ser	Gly	Ile	Ser	Thr	Thr	Phe	Pro	Lys	Leu	Arg	Met	Arg	Val	Arg	
		** :			245	•			•	250					255	_	
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;	Asn	Ile	Leu	Ile	His	Asn	Asn	Tvr	Lvs	Ser	Ala	Thr	His	Glu	Asn	Asn	
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. I	His	Ser	Val.	Cys	Leu	Pro	Ala	Ala	Thr	Gln	Asn	Ile	Pro	Pro	Gly	Ser	-3
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•	Thr	Ala	Tyr	Val	Thr	Gly	Trp	Gly	Ala.	_Gln	Glu	Tyr	Ala	Gly	His	Thr	
	305					310		•			315	-			•	320	3 *
7	7a1	Pro	Glu	Leu	Arg	Gln	Glv	Gln	Va1	Ara	Tle	Tle	Ser	Asn	Asp	Val	
					325		3			330					335		
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		37Ò					375					380					
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G	ly	Ile	Val	Ser,	Trp	Gly	Asp	Gln	Cys	Gly	Leu	Pro	Asp	Lys	Pro	Glv	
	85	-17				390					395		-	-		400	
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 tttttgatct gtatgttttt cattttcatt cagcaagttt ttttttttt tcagagtctt 600
 actotyttyc coagyotyga gracaytyyt gcaatotoay otoactycaa cototycoto 660
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375 () 380 😘 Lys Gln Ala Thr Ser Pro Ala Ser Lys Lys Pro Ala Gln Glu Gly Gly 390 (2015) 395 (2015) 400 (2015) 395 (2015) 400 (2015) 400 (2015) 400 (2015) 400 (2015) 400 (2015) 400 (2015) Lys Gly Gly Ser Glu Lys Pro Lys Arg Pro Val Ser Ala Met Phe Ile 405 Tym E418 - 410 Phe Ser Glu Glu Lys Arg Arg Gln Leu Gln Glu Glu Arg Pro Glu Leu 420 and 1 125 at 1 2 1 430 at 2 4 25 Ser Glu Ser Glu Leu Thr Arg Leu Leu Ala Arg Met Trp Asn Asp Leu aa 435 Ser Glu Lys Lys Lys Ala Lys Tyr Lys Ala Arg Glu Ala Ala Leu Lys Ala Gln Ser Glu Arg Lys Pro Gly Gly Glu Arg Glu Glu Arg Gly Lys . 470 Leu Pro Glu Ser Pro Lys Arg Ala Glu Glu Ile Trp Gln Gln Ser Val 495 490 ্ৰাক কেন্দ্ৰী ব Ile Gly Asp Tyr Leu Ala Arg Phe Lys Asn Asp Arg Val Lys Ala Leu 510 **510** 505 ். இது மாவிரு நிருக்கிரும் இரு நிருக்கி Lys Ala Met Glu Met Thr Trp Asn Asn Met Glu Lys Lys Glu Lys Leu ର ହଳ ବିଲିଲ୍ଲ , **515** ବିଲିଲ୍ଲ କିଲ୍ଲ କିଲ୍ଲ କିଥିଲି ଓ ଅଟେ ଓ ଅଟେ ଅଟେ ବିଲ୍ଲ <mark>(525</mark> ଅଟେ ସ୍ଥରୀ ଅଟେ କିଲ୍ଲ କିଥିଲେ । ବିଲ୍ଲାର କ୍ରେମ୍ବର ପ୍ରଥର ବିଲ୍ଲା ବିଲ୍ଲ କ୍ରେମ୍ବର କ୍ରେମ୍ବର ଅଟେ ଅଟେ ଅଟେ ଅଟେ ଅନ୍ୟର୍ଶ ଅଟେ କର୍ମ କର୍ମ ଅଟେ ଅଟେ ଅଟେ ଅଟେ ଅଟେ Met Trp Ile Lys Lys Ala Ala Glu Asp Gln Lys Arg Tyr Glu Arg Glu 284 (4.530), Provided to Section 535 provided to the Constant of the Constant Leu Ser Glu Met Arg Ala Pro Pro Ala Ala Thr Asn Ser Ser Lys Lys 555 550 Met Lys Phe Gln Gly Glu Pro Lys Lys Pro Pro Met Asn Gly Tyr Gln 570 565 Street Beerly Lys Phe Ser Gln Glu Leu Leu Ser Asn Gly Glu Leu Asn His Leu Pro 585 580 in the contract of the contrac Leu Lys Glu Arg Met Val Glu Ile Gly Ser Arg Trp Gln Arg Ile Ser 198 --อสุดกรุงกับ 595 มะ โดงการสุดให้ที่วัด 1600ใต้ ซึ่ง เการะก็กรก็ 605/กรณี เรื่องว่ากรก อาษารู้การสุดการ เพื่ออสุดเด็จสอย (อะสาร์สิทธิการ) เพื่อเกาะ เกาะ เกาะ เกียงให้ก็ก็การ เกาะ Gln Ser Gln Lys Glu His Tyr Lys Lys Leu Ala Glu Glu Gln Gln Lys 615 Gln Tyr Lys Val His Leu Asp Leu Trp Val Lys Ser Leu Ser Pro Gln Asp Arg Ala Ala Tyr Lys Glu Tyr Ile Ser Asn Lys Arg Lys Ser Met 650 the second Thr Lys Leu Arg Gly Pro Asn Pro Lys Ser Ser Arg Thr Thr Leu Gln 665

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Gly Asp Glu Asn Glu Glu Asp Asp Glu Asp Glu Asp Asp Asp Glu Asp
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Ser Asp Leu Leu Glu Trp Ile Arg Arg Thr Ile Pro Trp Leu Glu Asn
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stek<400>.81 million transferance groupe had by the signification of the contraction of 
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Gly Lys Val Ile Lys Cys Lys Ala Ala Val Leu Trp Glu Gln Lys Gln
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  Pro Phe Ser lle Glu Glu Ile Glu Val Ala Pro Pro Lys Thr Lys Glu 🦠 🤻
arendo promune<mark>r:35</mark>, postano esporteda o monta40, que en el arenda en el conf<mark>a</mark>5 sincipo e figurar e tipo
Per a cuerco e regular como promoto en el como porte el como en el
  Val Arg Ile Lys Ile Leu Ala Thr Gly Ile Cys Arg Thr Asp Asp His
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 Val Ile Lys Gly Thr Met Val Ser Lys Phe Pro Val Ile Val Gly His
 មិសី 65 ហា បទទី ។ មី មានប្រទេ70១ ខ្លួនមានមានមិន ១០១១.75 ។ ខេត្តសុខមិនមាន <mark>ទី៣៤ 80</mark>១៩៥៦៤
មិសី ឱានស្វាក់ ទី៣០១៨ ការស្វាយមាន ម៉ែន សង្គារពីមានការសង្គារការ មានបានសុខមាន បានស្វាស់ បានការប្រទេសម៉ូន ហ៊ុន
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                                                                                                                                  105
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                                                                                                                                                                                                               كبور كد
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Val	Phe 210	Gly	Leu	Arg	Gly	Val 215	Gly	Leu	Ser	Val	11e 220	Met	Gly	Суѕ	Lys	មេកភ មន្តិ៖
Ser 225	Ala	Gly	Ala	Ser	Arg 230	Ile	Ile	Gly	Ile	Asp 235	. Leu	Asn	Lys	Asp	Lys 240	A.
Phe	Glu	Lys	Ala	Met 245	Ala	Val	Gly	Ala	Thr. 250	Glu	Cys	Ile	Ser (0.	Pro 255	Lys	. 1 9 I - -
		•	260	Pro	•	. " .		265	72.7°		2.015		270	eler Galle		* 35.
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Asp	Ala 290		Ala	Ser	Cys	His 295	Met	Asn	Tyr	Gly	Thr 300	Ser	Val	Val	Val	ast E£
Gly 305	Val	Pro	Pro	Ser	Ala 310	Lys	Met	Leu	Thr	Tyr 315	Asp	Pro	Met	Leu	Leu 320	
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Asp		Asp 355	Gln	Leu.	Ile. Sec	Thr	His 360	Val,	Leu	Pro,	Phe	Lys 365	Lys	Ile	Ser	5 3 Q
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. J.s.				٠ ٠٠٠. 		135		i s tuvis	ing Singapan		140	ing. Singap			in the
	Leu			Gly							Gly	Thr	Thr	Tyr	Lys
145	i grand	s 11	٠.,٠.	1 12	150	5 0 4	\$ P			155		3 5		25	160
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	Leu	ini Sana		ere eger e La calaba	inger i e Lila a i g	4-11-5-29 4-11-5-1					•				2.00
i i daga ta	: 6:	gasti.				- 5 kg	jar v	4	i.		* 2			, Y	rajay wili di Karamatan
<210)> 94	1 .										•			4-1- 5-6
	l> 1(2> Pi							•							i sujit
-	3> Ho		sapi	ens							٠,				0 2747 1 3437
	•		_											* * 1. * 1. * 1. * 1. * 1. * 1. * 1. *	
	0 > 94		<i>c</i> 1	Ala	Thr	T.All	Gln	Hie	Glü	λla	Thr	Δla	Δla	Thr	T.eu
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		al The			303 °	ng yek		. 5				**	ing sold Sold was so		
		_	His	Ala	Asp	Ser	Val	Ala	Glu	Leu	Gly	Glu	Gln 30	Ile	Asp
	ina 75 ma		20		111					, · · ·:		··no		35-1	$(\mathbb{S}^n, I_{(1)}, \mathbb{S}^n)$
Asn	Leu	Gln	Arg	Val	Lys	Gln	Lys	Leu	Glu	Lys				Glu	Met
	Selection 1	. 35	. * *	, /, /	1.3	is the first of the second	40	* - <u>.</u> .		· .		45		· 14, 3	profite to
:: Lvs-	Met	Glu	Ile	-Asp	Asp	Leu	Ala	Cys	Asn	Met	Glu	Val	Ile	Ser	Lys
	50			-		55		, -			60				* N
	T	C1	3	Leu	Clu	Tree	Mat	Ove	ara	Thr	T.em	Glu	Δen	Gln	Val
65	гÀг	GTĀ	ASII	rea	70	пуз	MEC	Cys	ALG	75	Deu	GIU	nap	350	80
Ser	Gla	T.eu	Tare	Thr	Gln	Glu	Glu	Glu	Gln	Gln	Ara	Leu	İle	Asn	Glu
Jer	010	Dea	Lly 3	85		1,331	ķ¢ļ₩.	ΔF.	90	Y-3:	77.5	ŢĀĀ		95	
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Leu	Thr	Ala			ء راد د			48.	ا آھين	. E - S				1.5	7 -1 - 1
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	0> 99		4)			कृत्य	210	قه د ۱۰		24	*. *.		• • •	.	
	1> 99			. •				-	• '				-		
<213	2> PI 3> Ho	owo, a	sapie	ens 🤌		(a, 3*	y JJ		1	÷		٠.	٠.,	•; •.	1 2
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<400	0> 95	5_			r <u>=</u> 50 cm				دا جد		**-		` - 121		Ole I
Lys 1	Ile	Leu	Pro	Leu 5	Asn	Gly	Asn	Leu	Gln 10	Ala	val	GIu	Leu	15	GIU
Lys	Arg:	Thr	Ser 20	Ser	Leu	Arg	Ile	Lys 25	Met	Phe	Arg	Ala	Thr 30	Arg	Val
		•				•									

Thr Ser Thr Ser Arg Phe Leu Asn Pro Tyr Val Val Cys Phe Leu Val
35 40 45

Leu Pro Gly Val Val Ile Leu Ala Val Pro Ile Ala Leu Leu Val Tyr
50 55 60

Phe Leu Ala Phe Asp Gln Lys Ser Tyr Phe Tyr Trp Ser Asn Phe Pro 65 70 75 80

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Gly Ile Pro

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Pro Val Gly Glu Asp Ser Leu Val Ser Asp Arg Ser Lys Lys Glu Leu 35 40 45

Ser Pro Val Leu Thr Ser Glu Val His Ser Val Arg Ala Gly Arg His
50 55 60

Leu Ala Thr Lys Leu Asn Ile Leu Val Gln Gln His Phe Asp Leu Ala 65 70 75 80

Ser Thr Thr Ile Thr Asn Ile Pro Met Lys Glu Glu Gln His Ala Asn 85 90 95

Thr Ser Ala Asn Tyr Asp Val Glu Leu Leu His His Lys Asp Ala His 100 105 110

Val Asp Phe Leu Lys Ser Gly Asp Ser His Leu Gly Gly Gly Ser Arg 115 120 125

Glu Gly Ser Phe Lys Glu Thr Ile Thr Leu Lys Trp Cys Thr Pro Arg 130 135 140

Thr Asn Asn Ile Glu Leu His Tyr Cys Thr Gly Ala Tyr Arg Ile Ser 145 150 155 160

Pro Val Asp Val Asn Ser Arg Pro Ser Ser Cys Leu Thr Asn Phe Leu 165 170 175

2-1-1-2

Leu Asn Gly Arg Ser Val Leu Leu Glu Gln Pro Arg Lys Ser Gly Ser 180 185 190

Lys Val Ile Ser His Met Leu Ser Ser His Gly Gly Glu Ile Phe Leu 195 200 205

His Val Leu Ser Ser Ser Arg Ser Ile Leu Glu Asp Pro Pro Ser Ile 210 215 220

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Gly Glu Phe Met Arg Gly Lys Gln Ile Asn Ser Phe Ser Thr Pro Gln
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Arg Gly Lys Gly Glu Ile Thr Pro Ala Ala Tle Gln Lys Met Leu Asp 45

Asp Asn Asn His Leu Ile Gln Cys Ile Met Asp Ser Gln Asn Lys Gly
50 55 60

Lys Thr Ser Glu Cys Ser Gln Tyr Gln Gln Met Leu His Thr Asn Leu 65 70 75 80

Val Tyr Leu Ala Thr Ile Ala Asp Ser Asn Gln Asn Met Gln Ser Leu 85 90 95

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Asn Asn Glu Gln Leu Leu Val Asn Gln Gln Ala Ile Gln Ile Leu
35 40 45

Glu Lys Ile Ser Gln Pro Val Val Val Val Ala Ile Val Gly Leu Tyr
50 55 60

Arg Thr Gly Lys Ser Tyr Leu Met Asn His Leu Ala Gly Gln Asn His 65 70 75

Gly Phe Pro Leu Gly Ser Thr Val Glm Ser Glu Thr Lys Gly Ile Trp-

Met Trp Cys Val Pro His Pro Ser Lys Pro Asn His Thr Leu Val Leu

Leu Asp Thr Glu Gly Leu Gly Asp Val Glu Lys Gly Asp Pro Lys Asn 115

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Tyr Asn Ser Met Ser Thr Ile Asn His Gln Ala Leu Glu Gln Leu 145 150 155

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<213> Homo sapiens

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20
25
30

Lys Ile Ser Gln Pro Val Val Val Val Ala Ile Val Gly Leu Tyr Arg
35 40 45

Thr Gly Lys Ser Tyr Leu Met Asn His Leu Ala Gly Gln Asn His Gly
50 55 60

Phe Pro Leu Gly Ser Thr Val Gln Ser Glu Thr Lys Gly Ile Trp Met 65 70 75 80

Trp Cys Val Pro His Pro Ser Lys Pro Asn His Thr Leu Val Leu Leu

Asp Thr Glu Gly Leu Gly Asp Val Glu Lys Gly Asp Pro Lys Asn Asp 105 110

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Val Thr Asp

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<213> Homo sapiens

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Ile Asn Asp Pro Phe Ile Asp Leu Asn Tyr Met Val Tyr Met Phe Gln

Tyr Asp Ser Thr His Gly Lys Phe His Gly Thr Val Glu Ala Glu Asn _____50_______60---

Gly Lys Leu Val Ile Asn Gly Asn Pro Ile Thr Ile Phe Gln Glu Arg 65

Asp Pro Ser Lys Ile Lys Trp Gly Asp Ala Gly Ala Glu Tyr Val Val

Glu Ser Thr Gly Val Phe Thr Thr Met Glu Lys Ala Gly Ala His Leu 105

Gln Gly Gly Ala Lys Arg Val Ile Ile Ser Ala Pro 120

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Gln Ser Ala Ala Ser Ser Phe Ala Ser Pro Ala Glu Pro His Arg Ser 10 15 mm

<213> Homo sapiens

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                                                                                            25
     Gly Arg Leu Val Thr Arg Ala Ala Phe Asn Ser Gly Lys Val Asp Ile
                            35
                                                                             40
                                                                                                                                 45
                             ٠. _
    Val Ala Ile Asn Asp Pro Phe Ile Asp Leu Asn Tyr Met Val Tyr Met
                                       55
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    Phe Gln Tyr Asp Ser Thr His Gly Lys Phe His Gly Thr Val Glu Ala
Glu Asn Gly Lys Leu Val Ile Asn Gly Asn Pro Ile Thr Ile Phe Gln
                                                                                                      90.
    Glu Arg Asp Pro Ser Lys Ile Lys Trp Gly Asp Thr Gly Ala Glu Tyr
                                                                                         105
                                    100
                                                                                                                                             110
    Val Val Glu Ser Thr Gly Val Phe Thr Thr Met Glu Lys Ala Gly
                                                                              120
                                                                                                                                   125
    <212> DNA 6d a m Agreem | Depty | pour mark we have see
                                                                                                    je selj siletoski t
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    ggcggccgcg gcgggccagc ggcggagccg tgtagcggag aageteeece teeetgette 180
    cettggccga gccgggggc cgcgcacg cggccgtcca gagcgggctc cccacccctc 240
    gactectgeg accegeaceg caceeceace egggeeegga ggatgatgaa geteaagteg 300
    aaccagacco gcacctacga cggcgacggc tacaagaagc gggccgcatg cctgtgtttc 360
    cgcagcgaga gcgaggagga ggtgctactc gtgagcagta gtcgccatcc agacagatgg 420
    attgtccctg gaggaggcat ggagcccgag gaggagccaa gtgtggcagc agttcgtgaa 480
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ccaagtgcca aaaaaaggcc tgattaggcc ctgaaattca gtgaaattct gcctgaagaa 420
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  and other rest. The control of
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tgatagtgat agtgacagcg atgatgatga agatgatgtg catgtcacta taggagacat 420
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Тут	Glr	Met	Leu 20		Asn	Ası	ı Tr	o Gl: 25		Leu	Ser	Ser	Phe 30	_	g Gly
Glr	g] Glu	Phe 35	val	Trp	Asp	Тут	Va]		e Leu	Asp	Glu	Ala 45		Lys	Ile
Lys	Thr 50		Ser	Thr	Lys	Ser 55		Ile	e Cys	Ala	Arg 60		Ile	Pro	Ala
	Așn	Arg	Leu	Leu	Leu 70		Gly	Thr	Pro	Ile 75		Asn	Asn	Leu	Gln 80
Glu	Leu	Trp	Ser	Leu 85	Phe	- A sp	্Phe া	Ala	Cys 90		Gly	Ser	Leu	Leu 95	_
Thr	Leu	Lys	Thr 100	Phe	Lys	Met	Glu	Tyr 105		Asn	Pro	Ile	Thr 110	Arg	Ala
Arg	Glu	Lys 115	Asp	Ala	Thr	Pro	Gly 120		Lys	Ala	Leu	Gly 125	Phe	Lys	Ile
Ser	Glu 130		Leu	Met	Ala	Ile 135		Lys	Pro	Tyr	Phe 140	Leu	Arg	Arg	Thr
			Val									Glu			Leu 160
Asn			Asn					Ala		Cys		Met	Pro		Leu
Ser	Arg	Arg	Asn 180	Asp	Leu	Ile ∵⊖	Ile 맛가지	Trp 185	Ile	Arg	Leu :	V al	Pro	Leu	
Glu		Ile 195	Tyr						Leu						Leu
Leu			Thr								Gly 220	Val	Leu	Lys	Lys
Leu 225			His v3										٠Ĵ٠;	. :	
Asn			Thr				rej.		, 5s	· ^T	771 S.	•		٠.	
<211)> 11 l> 10 !> PR	7.	prop.	1 449 201	¥†£	च क्री इ.स.	ga i	28%	† . ',	4-,11	v H		कपूत्र"	, 5 %:	•21 [™] •
	> Ho	mo s	apie		ψχ.Υ 1+1		Gir.	· **,	1		. 54 . 1	es jin	+ Tr		· · · · · · · · · · · · · · · · · · ·
			Val	Ile :	Lys :	Asp	The	Lys	Leu	Leu	Cys	Tyr	Lys	Ser	Ser

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1				5			:		10		-			15	- `
Lys	Asp	Gln	G1n 20	Pro	Gln	Met	Glu	Leu -25	Pro	Leu	Gln		Cys 30	Asn	Ile
Thr	Tyr	Ile 35	Pro	Lys	Asp	Ser	Lys 40	Lys	Lys	Lys	His	Glu 45	Leu	Lys	Ile
Thr	Gln 50	Gln	Gly	Thr	Asp	Pro 55	Leu	Val	Leu	Ala	Val 60	Gln	Ser	Lys	Glu
Gln 65	Ala	G1ú	Gln	Trp	Leu 70		Val	Ile	Lys	Glu 75		Tyr	Ser	Gly	Cys 80
Ser	Gly	Pro	Val	Asp 85	Ser	Glu	Cys	Pro	Pro 90	Pro	Pro	Ser	Ser	Pro 95	Val
His	Lys	Ala	Glu 100	Leu	Glu	Lys	Lys	Leu 105		.Ser		स्टरी <u>अ</u> न्		1904.15	1 m 3 m
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Asp	Lys	Asn	Glu 20	Ile				25	Tyr	Arg	Tyr		30	Trp	Lys
Leu	Gly	Asp 35	Asp		Asp	Leu	Ile 40	Val	Arg	CAa	Glu	His 45	Asp	Gly	Val
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Leu	Asp	Ser	Gln	Arg 85		Ala	° Val	Ile	Ala • 90		Glű	Leu	Lys	95	Asn
Ser	Tyr	Lys	Leu 100	Ala	Arg	Trp	Thr	Cys 105	Cys	Ala		Lèu		Gly	Ser
Glu	Tyr	Leu 115		Leu	Gly	Tyr	Val 120	Ser	Arg	Tyr	His	Val 125	Lys	•	Ser
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Ala	Ser	Gln	Ile	Asn	Leu	Ser	Val	Glu	Asn	Alá		*	j. G	•	

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Ala	Glu	Ala	Leu	Arg	Ala	Pro	Arg	Ala	Gly	Gln	Pro	Leu	Gln	Leu	Leu
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His	Phe	Ala	Glu	Tyr 85	Ala	Gly	Arg	Leu	Gly 90	Val	Gly	Ala	Ala	Thr 95	His
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cacteegetg tegeccacec geatcaceeg getgeaggag aaggaggace tgeaggaget
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gegegeeege etgeagetgg agetgageaa agtgegtgaa gagtttaagg agetgaaage
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	ttgttgaatt caagatggaa					420
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_	aaacctc			MAG - A A		547
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	ggaagatece acagteteag					720
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	gctctactta cagaacagcc agcaataaag ctatgtctga					838
	ageacaaag clatyciya		-uccaaaaa	uuuuuaaada	addecada?	339
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	cgactggtac cagaagcagg	ggcctgggcc	-eteeegegae-	tacagecact	actacacgac	240
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Lys Glu Lys Asn Ile Lys Arg Gly Gly Asn Arg Phe Glu Pro Tyr Ser
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Asn Pro Thr Lys Arg Tyr Arg Ala Phe Ile Thr Asn Ile Pro Phe Asp
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                                                 70
Val Lys Trp Gln Ser Leu Lys Asp Leu Val Lys Glu Lys Val Gly Glu
                                                                                          90
                                       85
Val Thr Tyr Val Glu Leu Leu Met Asp Ala Glu Gly Lys Ser Arg Gly
                                                                               105
                             100
Cys Ala Val Val Glu Phe Lys Met Glu Glu Ser Met Lys Lys Ala Ala
                                                                    120
                   115
Glu Val Leu Asn Lys His Ser Leu Ser Gly Arg Pro Leu Lys Val Lys
130
                                                                                                            140
Glu Asp Pro Asp Gly Glu His Ala Arg Arg Ala Met Gln Lys Ala Gly
145
                                                                                                155
Arg Leu Gly Ser Thr Val Phe Val Ala Asn Leu Asp Tyr Lys Val Gly
                                       165
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Trp Lys Lys Leu Lys Glu Val Phe Ser Met Ala Gly Val Val Val Arg
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Ala Asp Ile Leu Glu Asp Lys Asp Gly Lys Ser Arg Gly Ile Gly Ile
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Val Thr Phe Glu Gln Ser Ile Glu Ala Val Gln Ala Ile Ser Met Phe
                                                                                          220
  210 215
Asn Gly Gln Leu Leu Phe Asp Arg Pro Met His Val Lys Met Asp Glu
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Met Gln Glu Ser Val Leu Asp Phe Asp Lys Pro Ser Ser Ala Ile Pro 345 340 Thr Ser Gln Pro Pro Ser Ala Thr Pro Gly Ser Pro Val Ala Ser Lys 360 355 · Glu Gln Asn Leu Ser Ser Gln Ser Asp Phe Leu Gln Glu Pro Leu Gln 370 375 380 Val Phe Asn Val Asn Ala Pro Leu Pro Pro Arg Lys Glu Gln Glu Ile 390 395 Lys Glu Ser Pro Tyr Ser Pro Gly Tyr Asn Gln Ser Phe Thr Thr Ala 410 405 Ser Thr Gln Thr Pro Pro Gln Cys Gln Leu Pro Ser Ile His Val Glu 420 425 Gln Thr Val His Ser Gln Glu Thr Ala Ala Asn Tyr His Pro Asp Gly 440 Thr Ile Gln Val Ser Asn Gly Ser Leu Ala Phe Tyr Pro Ala Gln Thr 455 460 Asn Val Phe Pro Arg Pro Thr Gln Pro Phe Val Asn Ser Arg Gly Ser 470 475 Val Arg Gly Cys Thr Arg Gly Gly Arg Leu Ile Thr Asn Ser Tyr Arg Ser Pro Gly Gly Tyr Lys Gly Phe Asp Thr Tyr Arg Gly Leu Pro Ser 500 505 Ile Ser Asn Gly Asn Tyr Ser Gln Leu Gln Phe Gln Ala Arg Glu Tyr 515 520 Ser Gly Ala Pro Tyr Ser Gln Arg Asp Asn Phe Gln Gln Cys Tyr Lys 540 535 ... Arg Gly Gly Thr Ser Gly Gly Pro Arg Ala Asn Ser Arg Ala Gly Trp **550** 555 Ser Asp Ser Ser Gln Val Ser Ser Pro Glu Arg Asp Asn Glu Thr Phe 575 565 570 Asn Ser Gly Asp Ser Gly Gln Gly Asp Ser Arg Ser Met Thr Pro Val 580 590 585 Asp Val Pro Val Thr Asn Pro Ala Ala Thr Ile Leu Pro Val His Val 595 605 Tyr Pro Leu Pro Gln Gln Met Arg Val Ala Phe Ser Ala Ala Arg Thr 610 615 Ser Asn Leu Ala Pro Gly Thr Leu Asp Gln Pro Ile Val Phe Asp Leu 625 630 635 640 Leu Leu Asn Asn Leu Gly Glu Thr Phe Asp Leu Gln Leu Gly Arg Phe 185 E 8 1 645 650 Asn Cys Pro Val Asn Gly Thr Tyr Val Phe Ile Phe His Met Leu Lys 660 665 Leu Ala Val Asn Val Pro Leu Tyr Val Asn Leu Met Lys Asn Glu Glu 675 680 685 Val Leu Val Ser Ala Tyr Ala Asn Asp Gly Ala Pro Asp His Glu Thr 690 700 . 695 Ala Ser Asn His Ala Ile Leu Gln Leu Phe Gln Gly Asp Gln Ile Trp 7.05 710 715 Leu Arg Leu His Arg Gly Ala Ile Tyr Gly Ser Ser Trp Lys Tyr Ser 725 730 Thr Phe Ser Gly Tyr Leu Leu Tyr Gln Asp 740

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410 405 Leu Gly Ile Trp Gly Glu Gly Thr Pro Phe Arg Glu Phe Ser Asp Phe 425 430 420 Ile Gln Ala Val Glu Arg Arg Gly Val Gly Ala Met Glu Ile Val Ala 440 445 Met Asp Met Lys Leu Arg Gly Met Tyr Ile Ala Arg Gln Leu Ser Phe 460 455 Thr Gly Val Thr Phe Lys Ile Glu Glu Val Leu Leu Ser Gln Ser Tyr 470 475 Val Lys Met Tyr Asn Lys Ala Val Lys Leu Trp Val Ile Ala Arg Glu 485 490 Arg Phe Gln Gln Ala Ala Asp Leu Ile Asp Ala Glu Gln Arg Met Lys 505 Lys Ser Met Trp Gly Gln Phe Trp Ser Ala His Gln Arg Phe Phe Lys 520 Tyr Leu Cys Ile Ala Ser Lys Val Lys Arg Val Val Gln Leu Ala Arg 535 Glu Glu Ile Lys Asn Gly Lys Cys Val Val Ile Gly Leu Gln Ser Thr 550 555 Gly Glu Ala Arg Thr Leu Glu Ala Leu Glu Glu Gly Gly Gly Glu Leu 570 565 Asn Asp Phe Val Ser Thr Ala Lys Gly Val Leu Gln Ser Leu Ile Glu 585 580 Lys His Phe Pro Ala Pro Asp Arg Lys Lys Leu Tyr Ser Leu Leu Gly 600 Ile Asp Leu Thr Ala Pro Ser Asn Asn Ser Ser Pro Arg Asp Ser Pro 615 620 Cys Lys Glu Asn Lys Ile Lys Lys Arg Lys Gly Glu Glu Ile Thr Arg 635 630 Glu Ala Lys Lys Ala Arg Lys Val Gly Gly Leu Thr Gly Ser Ser Ser - **65**0 645 Asp Asp Ser Gly Ser Glu Ser Asp Ala Ser Asp Asn Glu Glu Ser Asp 660 665670 Tyr Glu Ser Ser Lys Asn Met Ser Ser Gly Asp Asp Asp Phe Asn • 680 685 Pro Phe Leu Asp Glu Ser Asn Glu Asp Asp Glu Asn Asp Pro Trp Leu 700 695 Ile 705 (数1.6.35 12.16) <210> 187 <211> 595 <212> PRT <213> Homo sapien Carrier green to South Soft <400> 187 Glu Ser Pro Arg His Arg Gly Glu Gly Gly Glu Trp Gly Pro Gly 10 g / 4 g / g / Six -15 g / 60

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Glu	Trp	Glý	Pro	Ser	Pro	Ser	Gly	His	Gly	Asp	Gly	Pro	Arq	Arq	Arq
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Pro	Arg	Lys	Arg	Arg	Gly	Arg	Lys	Gly	Arq	Met	GÍV	Ara	Gln	His	Glu
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Ala	Ala	Ala	Thr	Ala	Ala	Thr	Ala	Ala	Thr	Ala	Thr	Glv	Glv	Thr	Ala
	210					215			-	7.3	220	3	3		
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Arg	Gly	Arq	Ala	Arg	Glv	Pro	Arg	Gln	Gln	Glv	Ara	Ara	Ara	His	
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Asp	Ala	Thr	Thr	Ile	Leu	Gly	Leu	Gly	Thr	Pro	Ser	Glv	Glu	Gln	Arg
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Ala	Asp	Gln	Ser	Gln	Ala	Leu	Pro	Ala	Leu	Ala	Glv		Ala	Ala	Ala
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Gly	Arg	Gly	Arg	Arg	Gly	Gly	Trp	Arg	Gly	Gly	Arq	Arg	Gly	Gly	Ser
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Ala	Gly.				-						Gly	Arq	Gly	Arg	Gly
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Àrg	Arg	Gly	Arg	Gly	Pro	Pro	Ala	Ala	Gly	Ala	Ala	Gln	Val	Ser	Ala
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Arg	Gly	Arg	Arg	Ala	Arg	Gly	Gln	Arq	Ala	Gly	Glu	Glu	Ala	Gln	Asp
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Gly	Leu	Leu	Pro	Arg	Glv	Arg	Asp	Ara			Leu	Ara	Pro		Asp
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Arg Pro Gly Pro Arg Arg Pro Ala Arg Arg Pro Arg Gly Glu Leu Ile 520 - Jan 525 - 1 Pro Arg Arg Pro Asp Pro Ala Ala Pro Ser Glu Glu Gly Leu Arg Met 535 540 Glu Ser Ser Val Asp Asp Gly Ala Thr Ala Thr Thr Ala Asp Ala Ala 560 grading 555 grading at 1996 560 550 Ser Gly Glu Ala Pro Glu Ala Gly Pro Ser Pro Ser His Ser Pro Thr 570 565 Met Cys Gln Thr Gly Gly Pro Gly Pro Pro Pro Pro Gln Pro Pro Arg 585 Frankling 1 1 590 H Trp Leu Pro 595 <210> 188 <211> 376 <212> PRT <213> Homo sapien <400> 188 中文化 经存货格 如何 数据 生物 人名德人尔特 Glu Met Arg Lys Phe Asp Val Pro Ser Met Glu Ser Thr Leu Asn Gln Pro Ala Met Leu Glu Thr Leu Tyr Ser Asp Pro His Tyr Arg Ala His Phe Pro Asn Pro Arg Pro Asp Thr Asn Lys Asp Val Tyr Lys Val Leu 1.40 (1.10) par with Se*4546 for the company Pro Glu Ser Lys Lys Ala Pro Gly Ser Gly Ala Val Phe Glu Arg Asn Gly Pro His Ala Ser Ser Ser Gly Val Leu Pro Leu Gly Leu Gln Pro Ala Pro Gly Leu Ser Lys Ser Leu Ser Ser Gln Val Trp Gln Pro Ser ्रहरू**90**% इंड वृद्धक केला र मार्थ **95**ला ुर्द्धकी 85 Pro Asp Pro Trp His Pro Gly Glu Gln Ser Cys Glu Leu Ser Thr Cys 7 J105 7 Page 455 708 200 110 100 Arg Gln Gln Leu Glu Leu Ile Arg Leu Gln Met Glu Gln Met Gln Leu .120_{00 ave} the put allowed the company Gln Asn Gly Ala Met Cys His His Pro Ala Ala Phe Ala Pro Leu Leu (p. 1966) gett reg**140**位 (396) (115) 1 (146) 135 Pro Thr Leu Glu Pro Ala Gln Trp Leu Ser Ile Leu Asn Ser Asn Glu 150 155 His Leu Leu Lys Glu Lys Glu Leu Leu Ile Asp Lys Gln Arg Lys His 170 175 165 Ile Ser Gln Leu Glu Gln Lys Val Arg Glu Ser Glu Leu Gln Val His 190 L 185 . Ser Ala Leu Leu Gly Arg Pro Ala Pro Phe Gly Asp Val Cys Leu Leu 205 200 195 Arg Leu Gln Glu Leu Gln Arg Glu Asn Thr Phe Leu Arg Ala Gln Phe 215 220 Ala Gln Lys Thr Glu Ala Leu Ser Lys Glu Lys Met Glu Leu Glu Lys 230 235 Lys Leu Ser Ala Ser Glu Val Glu Ile Gln Leu Ile Arg Glu Ser Leu 245 250 Lys Val Thr Leu Gln Lys His Ser Glu Glu Gly Lys Lys Gln Glu Glu 265 Arg Val Lys Gly Arg Asp Lys His Ile Asn Asn Leu Lys Lys Cys

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Gln Lys Glu Ser Glu Gln Asn Arg Glu Lys Gln Gln Arg Ile GTu Thr
   290 295
                                  300
Leu Glu Arg Tyr Leu Ala Asp Leu Pro Thr Leu Glu Asp His Gln Lys
305 310
                              315
Gln Thr Glu Gln Leu Lys Asp Ala Glu Leu Lys Asn Thr Glu Leu Gln
                            330 335
     325 a 4 325
Glu Arg Val Ala Glu Leu Glu Thr Leu Leu Glu Asp Thr Gln Ala Thr
                         345 350
    340
Cys Arg Glu Lys Glu Val Gln Leu Glu Ser Leu Arg Gln Arg Glu Ala
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Asp Leu Ser Ser Ala Arg His Arg
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     <213> Homo sapien
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Thr Thr Ala Asn Arg Thr Gln Ser Leu Asn Tyr Gly Cys Ile Val Glu
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Asn Pro Gln Thr His Glu Val Leu His Tyr Val Glu Lys Pro Ser Thr
40 45
Phe Ile Ser Asp Ile Ile Asn Cys Gly Ile Tyr Leu Phe Ser Pro Glu
50 55
Ala Leu Lys Pro Leu Arg Asp Val Phe Gln Arg Asn Gln Gln Asp Gly
65, at 1, a Table 4 (2) in 70 The Theory of 1 2 75 The
Gln Leu Glu Asp Ser Pro Gly Leu Trp Pro Gly Ala Gly Thr Ile Arg
90
95
Leu Glu Gln Asp Val Phe Ser Ala Leu Ala Gly Gln Gly Gln Ile Tyr.
100 105
Val His Leu Thr Asp Gly Ile Trp Ser Gln Ile Lys Ser Ala Gly Ser
 125
Ala Leu Tyr Ala Ser Arg Leu Tyr Leu Ser Arg Tyr Gln Asp Thr His
 Pro Glu Arg Leu Ala Lys His Thr Pro Gly Gly Pro Trp Ile Arg Gly
(A) <210<sub>≥</sub> 190 (A) (B) (B)
    <211> 146
   <212> PRT
    <213> Homo sapien
    and the same of the same of the
    <400> 190: 38
Met Asp Pro Arg Ala Ser Leu Leu Leu Gly Asn Val Tyr Ile His
.1 5
                            10
                                           15
Pro Thr Ala Lys Val Ala Pro Ser Ala Val Leu Gly Pro Asn Val Ser
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1 5 10 15

Pro Thr Ala Lys Val Ala Pro Ser Ala Val Leu Gly Pro Asn Val Ser
20 25 30

Ile Gly Lys Gly Val Thr Val Gly Glu Gly Val Arg Leu Arg Glu Ser
35 40 45

Ile Val Leu His Gly Ala Thr Leu Gln Glu His Thr Cys Val Leu His
50 55 60

Ser Ile Val Gly Trp Gly Ser Thr Val Gly Arg Trp Ala Arg Val Glu

65 Gly Thr Pro Ser Asp Pro Asn Pro Asn Asp Pro Arg Ala Arg Met Asp 85 90 Ser Glu Ser Leu Phe Lys Asp Gly Lys Leu Leu Pro Ala Ile Thr Ile 100 105 1. Page 1. 1. 1. 1. Leu Gly Cys Arg Val Arg Ile Pro Ala Glu Val Leu Ile Leu Asn Ser 115 120 125 Ile Val Leu Pro His Lys Glu Leu Ser Arg Ser Phe Thr Asn Gln Ile 130 135 Ile Leu 145 (1933) 245 E <210> 191 <211> 704 <212> PRT <213> Homo sapien i kirin era i dari tak 1704 <400> 191 京都 化铁 and set 1000 1000 1000 Glu Gly Gly Cys Ala Ala Gly Arg Gly Arg Glu Leu Glu Pro Glu Leu Glu Pro Gly Pro Gly Ser Ala Leu Glu Pro Gly Glu Phe 5 20 25 317 448 844 330 年 學院 Glu Ile Val Asp Arg Ser Gln Leu Pro Gly Pro Gly Asp Leu Arg Ser 40 Ala Thr Arg Pro Arg Ala Ala Glu Gly Trp Ser Ala Pro Ile Leu Thr The state of the s 55 Leu Ala Arg Arg Ala Thr Gly Asn Leu Ser Ala Ser Cys Gly Ser Ala 75 Leu Arg Ala Ala Gly Leu Gly Gly Gly Asp Ser Gly Asp Gly Thr 85 90 Ala Arg Ala Ala Ser Lys Cys Gln Met Met Glu Glu Arg Ala Asn Leu 105 110 Met His Met Met Lys Leu Ser Ile Lys Val Leu Leu Gln Ser Ala Leu 120 Apr. 125 % This was Ser Leu Gly Arg Ser Leu Asp Ala Asp His Ala Pro Leu Gln Gln Phe 135 140 V. 1400 15 Phe Val Val Met Glu His Cys Leu Lys His Gly Leu Lys Val Lys Lys 4 d 150 155 Ser Phe Ile Gly Gln Asn Lys Ser Phe Phe Gly Pro Leu Glu Leu Val 165 170 - 175 - 175 - 175 - 175 Glu Lys Leu Cys Pro Glu Ala Ser Asp Ile Ala Thr Ser Val Arg Asn 180 185 190 Leu Pro Glu Leu Lys Thr Ala Val Gly Arg Gly Arg Ala Trp Leu Tyr 195 ~ (} . . 200 205 Leu Ala Leu Met Gln Lys Lys Leu Ala Asp Tyr Leu Lys Val Leu Ile 215 220 Asp Asn Lys His Leu Leu Ser Glu Phe Tyr Glu Pro Glu Ala Leu Met -230 235 Met Glu Glu Gly Met Val Ile Val Gly Leu Leu Val Gly Leu Asn 245 250 255 Val Leu Asp Ala Asn Leu Cys Leu Lys Gly Glu Asp Leu Asp Ser Gln 265 Val Gly Val Ile Asp Phe Ser Leu Tyr Leu Lys Asp Val Gln Asp Leu 280 285 Asp Gly Gly Lys Glu His Glu Arg Ile Thr Asp Val Leu Asp Gln Lys

295 300 Asn Tyr Val Glu Glu Leu Asn Arg His Leu Ser Cys Thr Val Gly Asp 310 315 Leu Gln Thr Lys Ile Asp Gly Leu Glu Lys Thr Asn Ser Lys Leu Gln 325 330 Glu Glu Leu Ser Ala Ala Thr Asp Arg Ile Cys Ser Leu Gln Glu Glu 340 345 Gln Gln Gln Leu Arg Glu Gln Asn Glu Leu Ile Arg Glu Arg Ser Glu 360 Lys Ser Val Glu Ile Thr Lys Gln Asp Thr Lys Val Glu Leu Glu Thr 375 380 Tyr Lys Gln Thr Arg Gln Gly Leu Asp Glu Met Tyr Ser Asp Val Trp 390 395 Lys Gln Leu Lys Glu Glu Lys Lys Val Arg Leu Glu Leu Glu Lys Glu 405 410 Leu Glu Leu Gln Ile Gly Met Lys Thr Glu Met Glu Ile Ala Met Lys 420 425 430 Leu Leu Glu Lys Asp Thr His Glu Lys Gln Asp Thr Leu Val Ala Leu
435 440 Arg Gln Gln Leu Glu Glu Val Lys Ala Ile Asn Leu Gln Met Phe His 450 455 Lys Ala Gin Asn Ala Glu Ser Ser Leu Gln Gln Lys Asn Glu Ala Ile 470 475 Thr Ser Phe Glu Gly Lys Thr Asn Gln Val Met Ser Ser Met Lys Gln 485 490 Met Glu Glu Arg Leu Gln His Ser Glu Arg Ala Arg Gln Gly Ala Glu 500 505 Glu Arg Ser His Lys Leu Gln Gln Glu Leu Gly Gly Arg Ile Gly Ala 515 520 525 Leu Gln Leu Gln Leu Ser Gln Leu His Glu Gln Cys Ser Ser Leu Glu 535 540 Lys Glu Leu Lys Ser Glu Lys Glu Gln Arg Gln Ala Leu Gln Arg Glu 550 Leu Gln His Glu Lys Asp Thr Ser Ser Leu Leu Arg Met Glu Leu Gln 565 570 Gln Val Glu Gly Leu Lys Lys Glu Leu Arg Glu Leu Gln Asp Glu Lys √ 580 ^{- ⊕ (} 585 Ala Glu Leu Gln Lys Ile Cys Glu Glu Gln Glu Gln Ala Leu Gln Glu 600 Met Gly Leu His Leu Ser Gln Ser Lys Leu Lys Met Glu Asp Ile Lys 615 620 Glu Val Asn Gln Ala Leu Lys Gly His Ala Trp Leu Lys Asp Asp Glu 630 635 Ala Thr His Cys Arg Gln Cys Glu Lys Glu Phe Ser Ile Ser Arg Arg 645 650 Lys His His Cys Arg Asn Cys Gly His Ile Phe Cys Asn Thr Cys Ser 660 665 Ser Asn Glu Leu Ala Leu Pro Ser Tyr Pro Lys Pro Val Arg Val Cys 680 Asp Ser Cys His Thr Leu Leu Gln Arg Cys Ser Ser Thr Ala Ser

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<211> 331

<212> PRT

<213> Homo sapien

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<213> Homo sapien

Lys Asn Ser Pro Leu Leu Ser Val Ser Ser Gln Thr Ile Thr Lys Glu 10 Asn Asn Arg Asn Val His Leu Glu His Ser Glu Gln Asn Pro Gly Ser

			20					25					30	· - .	
Ser	Ala	Gly 35	Asp	Thr	Ser	Ala	Ala 40	His	Gln	Val	Val	Leu 45	Gly	Glu	Asn
	50					55		_	_		60				Asp
65			•		70					75	_				Leu 80
		٠.		85			Thr		90		_			95	
			100				Asn	105				_	110		
		115		×			Asn 120		. 0		.•	125			·
*	130				***	135	Gly			. : -	140				
145					150	- •	Ser			155	4.7	52 1		•	160
		• .*		165			Ala		170			1 . 4.		175	15.44
			180				Lys	185				_	190		
		195	٠.				Leu 200					205			
1.	210					215	Gln				-220		,		i e
225	٠.				230		Leu Lys	,		235				•	240
		. :		245					250					255	
			260			٠.	Lys	265				-	270		. 413
	·	275	··· • • • • • •				Glu 280				_	285	e. See	F %	i. Est
	290		•			295	Asn			**	300	٠.		-	
305	,	•	• •		310	1	Leu			315			**		320
		. 1		325			Lys Leu		330	•			; •	335	T. Hegit
•			340				Val	345	7,00				350		
		355		٠.			360		•		_	365,		ne e. No sometal	
1.	370				÷	375					380	-	535.7		Leu
385		•			390	18.3	Thr		4	395					400
				405			Val		410				e de la companya de l	415	
			420					425					430		Met .
•		435	•	•			Leu 440					445		A.	
Leu	Ser 450	Gln	Pro	Ser	Gln	Pro 455	Ser	Ser	Pro	Leu	Pro 460	Gly	Ser	His	Gly

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Pro Phe Ala Glu Arg Thr Arg Leu Val Leu Lys Ala Lys Gly Ile Arg
                       40
                           45
His Glu Val Ile Asn Ile Asn Leu Lys Asn Lys Pro Glu Trp Phe Phe
Lys Lys Asn Pro Phe Gly Leu Val Pro Val Leu Glu Asn Ser Gln Gly
                                70
Gln Leu Ile Tyr Glu Ser Ala Ile Thr Cys Glu Tyr Leu Asp Glu Ala
             85 -
                           Tyr Pro Gly Lys Lys Leu Leu Pro Asp Asp Pro Tyr Glu Lys Ala Cys
                                     1. 1. 1. 110 3 W. T.
                          105
         100
Gln Lys Met Ile Leu Glu Leu Phe Ser Lys Val Pro Ser Leu Val Gly
115 120 125
Ser Phe Ile Arg Ser Gln Asn Lys Glu Asp Tyr Ala Gly Leu Lys Glu
                                    , 140 , som , su sav
                    135
Glu Phe Arg Lys Glu Phe Thr Lys Leu Glu Glu Val Leu Thr Asn Lys
                                155
                150
Lys Thr Thr Phe Phe Gly Gly Asn Ser Ile Ser Met Ile Asp Tyr Leu
                             170 : 💎 👾 🛬 🖯 😉 🛪 175% 🕹
             165
Ile Trp Pro Trp Phe Glu Arg Leu Glu Ala Met Lys Leu Asn Glu Cys
                          185 - 190
Val Asp His Thr Pro Lys Leu Lys Leu Trp Met Ala Ala Met Lys Glu
                      200 g205 (418)
      195
Asp Pro Thr Val Ser Ala Leu Leu Thr Ser Glu Lys Asp Trp Gln Gly
                   215
Phe Leu Glu Leu Tyr Leu Gln Asn Ser Pro Glu Ala Cys Asp Tyr Gly
                                        240
                230
                                 235
Leu
                 on was the first
    <210> 195
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                 1157 (41) S. S. S.
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Gln Thr Lys Ile Leu Glu Glu Asp Leu Glu Gln Ile Lys Leu Ser Leu
          Arg Glu Arg Gly Arg Glu Leu Thr Thr Gln Arg Gln Leu Met Gln Glu
             25 30.
          20
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Arg Ala Glu Glu Gly Lys Gly Pro Ser Lys Ala Gln Arg Gly Ser Leu

Glu His Met Lys Leu Ile Leu Arg Asp Lys Glu Lys Glu Val Glu Cys

40 3 3 3 3 4 5 3 3 4 7 6

55 60 Gln Gln Glu His Ile His Glu Leu Gln Glu Leu Lys Asp Gln Leu Glu 70 75 Gln Gln Leu Gln Gly Leu His Arg Lys Val Gly Glu Thr Ser Leu Leu 90 · 85 Leu Ser Gln Arg Glu Gln Glu Ile Val Val Leu Gln Gln Gln Leu Gln 105 100 Glu Ala Arg Glu Gln Gly Glu Leu Lys Glu Gln Ser Leu Gln Ser Gln 120 125 A Leu Asp Glu Ala Gln Arg Ala Leu Ala Gln The Mark Street Control 25 <210> 196 The state of the s <211> 102 <212> PRT <213> Homo sapien <400> 196 Met Ser Lys Arg Lys Ala Pro Gln Glu Thr Leu Asn Gly Gly Ile Thr 5 -10 Asp Met Leu Thr Glu Leu Ala Asn Phe Glu Lys Asn Val Ser Gln Ala 20 25 30 Ile His Lys Tyr Asn Ala Tyr Arg Lys Ala Ala Ser Val Ile Ala Lys appy (1886 - **35**일 - 전략 관계 원칙 40 45 Tyr Pro His Lys Ile Lys Ser Gly Ala Glu Ala Lys Lys Leu Pro Gly ੂਜ਼ੀ ਅ**50**ਾਰਕੀ ਅਹਿੰਦ ਨੂੰ ਉੱਤੇ ਤੋਂ ਹੈ 60 · 04 Val Gly Thr Lys Ile Ala Glu Lys Ile Asp Glu Phe Leu Ala Thr Gly
65 70 75 80 .75 Lys Leu Arg Lys Leu Glu Lys Ile Arg Gln Asp Asp Thr Ser Ser Ser 85 🚟 🦠 90 Ile Asn Phe Leu Thr Arg 100 <210> 197 <211> 138 <212> PRT <213> Homo sapien Glu Ala Asn Glu Val Thr Asp Ser Ala Tyr Met Gly Ser Glu Ser Thr 10 Tyr Ser Glu Cys Glu Thr Phe Thr Asp Glu Asp Thr Ser Thr Leu Val 20 30 25 His Pro Glu Leu Gln Pro Glu Gly Asp Ala Asp Ser Ala Gly Gly Ser _g.**45**m, 5.01€± 40 Ala Val Pro Ser Glu Cys Leu Asp Ala Met Glu Glu Pro Asp His Gly 60 Ext 32000 55⁻ Ala Leu Leu Leu Pro Gly Arg Pro His Pro His Gly Gln Ser Val 75 Ile Thr Val Ile Gly Gly Glu Glu His Phe Glu Asp Tyr Gly Glu Gly **30. 1 20. 19. 85**名 8元 (1971年) 90 Ser Glu Ala Glu Leu Ser Pro Glu Thr Leu Cys Asn Gly Gln Leu Gly

Cys Ser Asp Pro Ala Phe Leu Thr Pro Ser Pro Thr Lys Arg Leu Ser

125

100

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Lys Phe Met Cys Glu Val Gln Val Glu Gly Tyr Asn Tyr Thr Gly Met
       35 ..
Gly Asn Ser Thr Asn Lys Lys Asp Ala Gln Ser Asn Ala Ala Arg Asp
                      55 .
Phe Val Asn Tyr Leu Val Arg Ile Asn Glu Ile Lys Ser Glu Glu Val
65 70 75 80
Pro Ala Phe Gly Val Ala Ser Pro Pro Pro Leu Thr Asp Thr Pro Asp
 5.7 4.7 5 7 5. 185 5 7 1 1 4 4 4 4 5 9 0 5 8 6 6 7
Thr Thr Ala Asn
100 (
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CHA BAK 212> PRT BY THE BOTH BOTH
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                                10
Ala Thr Gln Glu Glu Leu Lys Lys Ala Tyr Arg Lys Leu Ala Leu Lys
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                             25
Tyr His Pro Asp Lys Asn Pro Asn Glu Gly Glu Lys Phe Lys Gln Ile
                         40
Ser Gln Ala Tyr Glu Val Leu Ser Asp Ala Lys Lys Arg Glu Leu Tyr
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Asp Lys Gly Gly Glu Gln Ala Ile Lys Glu Gly Gly Ala Gly Gly Gly
                  70
Phe Gly Ser Pro Met Asp Ile Phe Asp Met Phe Phe Gly Gly Gly
901
Arg Met Gln Arg Glu Arg Gly Lys Asn Val Val His Gln Leu Ser
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Val Thr Leu Glu Asp Leu Tyr Asn Gly Ala Thr Arg Lys Leu Ala
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                                           1986 1.24 DES
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                        10
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Glu Leu Lys Glu Leu Ile Gln Lys Glu Leu Thr Ile Gly Ser Lys Leu 35 40 45

Gln Asp Ala Glu Ile Ala Arg Leu Met Glu Asp Leu Asp Arg Asn Lys 50 55 60

Asp Gln Glu Val Asn Phe Gln Glu Tyr Val Thr Phe Leu Gly Ala Leu 65 70 75 80

Ala Leu Ile Tyr Asn Glu Ala Leu Lys Gly

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<213> Homo sapien

 Ado > 201

 Met Glu Thr Pro Ser Gln Arg Arg Ala Thr Arg Ser Gly Ala Gln Ala

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 Ser Ser Thr Pro Leu Ser Pro Thr Arg Ile Thr Arg Leu Gln Glu Lys

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 25

 30

 Glu Asp Leu Gln Glu Leu Asn Asp Arg Leu Ala Val Tyr Ile Asp Arg

 40
 45

 Val Arg Ser Leu Glu Thr Glu Asn Ala Gly Leu Arg Leu Arg Ile Thr

 50
 55

 60

 Glu Ser Glu Glu Val Val Ser Arg Glu Val Ser Gly Ile Lys Ala Ala

 65
 70

 75
 80

 Tyr Glu Ala Glu Leu Gly Asp Ala Arg Lys Thr Leu Asp Ser Val Ala

 85
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 Lys Glu Arg Ala Arg Leu Gln Leu Glu Leu Ser Lys Val Arg Glu Glu

 100
 105

 110

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Lys Glu Asp Pro Asp Gly Glu His Ala Arg Arg Ala Met Gln Lys Ala
145 150 155 160

Gly Arg Leu Gly Ser Thr Val Phe Val Ala Asn Leu Asp Tyr Lys Val
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<400> 203

Met Arg Leu Ala Val Gly Ala Leu Leu Val Cys Ala Val Leu Gly Leu 1 5 15 Cys Leu Ala Val Pro Asp Lys Thr Val Arg Trp Cys Ala Val Ser Glu 20 . . . 25 His Glu Ala Thr Lys Cys Gln Ser Phe Arg Asp His Met Lys Ser Val 40 IIe Pro Ser Asp Gly Pro Ser Val Ala Cys Val Lys Lys Ala Ser Tyr 55 Leu Asp Cys Ile Arg Ala Ile Ala Ala Asn Glu Ala Asp Ala Val Thr 70 75 garage Leu Asp Ala Gly Leu Val Tyr Asp Ala Tyr Leu Ala Pro Asn Asn Leu 90 2.95 Lys Pro Val Val Ala Glu Phe Tyr Gly Ser Lys Glu Asp Pro Gln Thr 100 105 Phe Tyr Tyr Ala Val Ala Val Val Lys Lys Asp Ser Gly Phe Gln Met 115 120 125 Asn Gln Leu Arg Gly Lys Lys Ser Cys His Thr Gly Leu Gly Arg Ser 135 Ala Gly Trp Asn Ile Pro Ile Gly Leu Leu Tyr Cys Asp Leu Pro Glu 150 325 155 Pro Arg Lys Pro

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<213> Homo sapien

<400> 205

Met Gln Ile Phe Val Lys Thr Leu Thr Gly Lys Thr Ile Thr Leu Glu 10 Val Glu Pro Ser Asp Thr Ile Glu Asn Val Lys Ala Lys Ile Gln Asp _____20 25 Lys Glu Gly Ile Pro Pro Asp Gln Gln Arg Leu Ile Phe Ala Gly Lys 40 Gln Leu Glu Asp Gly Arg Thr Leu Ser Asp Tyr Asn Ile Gln Lys Glu 55 Ser Thr Leu His Leu Val Leu Arg Leu Arg Gly Gly Met Gln Ile Phe 70 75 Val Lys Thr Leu Thr Gly Lys Thr Ile Thr Leu Glu Val Glu Pro Ser 90 Asp Thr Ile Glu Asn Val Lys Ala Lys Ile Gln Asp Lys Glu Gly Ile 100 105 Pro Pro Asp Gln Gln Arg Leu Ile Phe Ala Gly Lys Gln Leu Glu Asp 120 125 -Gly Arg Thr Leu Ser Asp Tyr Asn Ile Gln Lys Glu Ser Thr Leu His 130 Leu Val Leu Arg Leu Arg Gly Gly Met Gln Ile Phe Val Lys Thr Leu 155 150

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. <212> PRT

<213> Homo sapien

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<210> 207

<211> 175

<212> PRT

<213> Homo sapien

<400> 207

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165 170 175

<210> 208 <211> 177 <212> PRT <213> Homo sapien

<400> 208

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  Glu Arg Ser Ile Val Asp Tyr Lys Pro Asn Leu Asp Leu Leu Glu Gln
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  Gln His Gln Leu Ile Gln Glu Ala Leu Ile Phe Asp Asn Lys His Thr
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 Asn Glu Glu Ala Thr Gly Gln Phe His Val Tyr Pro Glu Leu Pro Lys
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 Pro Ser Ile Ser Ser Asn Asn Ser Asn Pro Val Glu Asp Lys Asp Ala
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 Val Ala Phe Thr Cys Glu Pro Glu Val Gln Asn Thr Thr Tyr Leu Trp
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<210> 213 <211> 142 <212> PRT <213> Homo sapien

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<210> 214 <211> 129 <212> PRT

<213> Homo sapien

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Arg	Pro 210	Gln	Pro	Pro	Pro	Pro 215		Leu	Pro	Pro	Pro 220	Pro	Glu	Ala	Gln
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Gly	Pro 370						_	Pro			Val 380		Val:	Ser	Ser
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Cys Phe Ser Arg Phe Ser Val Ser Pro Ala 485

Cys Phe Ser Arg Phe Ser Val Ser Pro Ala Leu Glu Thr Pro Gly Pro
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Pro Ala Arg Ala Pro Asp Ala Arg Pro Ala Gly Pro Val Glu Asn
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INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

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(21) International Application Number: PCT/US: (22) International Filing Date: 26 January 1999 (20) (30) Priority Data:	26.01.9 T T T	BY, CA, CH, CN, CU, CZ, DB, DK, EE, ES, FI, GB, GE, GH, GM, HR, HU, ID, IL, IS, IP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Burasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG). Published
(71) Applicant: CORIXA CORPORATION [US/US]; St 1124 Columbia Street, Scattle, WA 98104 (US).	uite 20	With international search report. Before the expiration of the time limit for amending the claim and to be republished in the event of the receipt of amendments.
(72) Inventors: REED, Steven, G.; 2843 – 122nd Pla Bellevue, WA 98005 (US). LODES, Michael, J. 36th Avenue S.W., Seattle, WA 98126 (US). FRU Tony, N.; P.O. Box 99232, Seattle, WA 99232-02 MOHAMATH, Raodoh; 4205 South Morgan, Sea 98118 (US).	; 9223 JDAKI 32 (US	9 December 1999 (09.12.99)
(74) Agents: MAKI, David, J. et al.; Seed and Ber 6300 Columbia Center, 701 Fifth Avenue, Seat 98104-7092 (US).	ту LL dle, W	À

(57) Abstract

Compounds and methods for treating lung cancer are provided. The inventive compounds include polypeptides containing at least a portion of a lung tumor protein. Vaccines and pharmaceutical compositions for immunotherapy of lung cancer comprising such polypeptides, or polynucleotides encoding such polypeptides, are also provided, together with polynucleotides for preparing the inventive polypeptides.

INTERNATIONAL SEARCH REPORT

Inte Sonal Application No PCT/US 99/01642

A CLASSIFICATION OF SUBJECT MATTER
1PC 6 C12N15/12 A611 C07K16/18 A61K35/14 C07K14/47 A61K38/17 According to International Patent Classification (IPC) or to both national classification and IPC Minimum documentation searched (classification system followed by classification symbols) IPC 6 C12N C12Q A61K C07K Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Electronic data base consulted during the International search (name of data base and, where practical, search terms used) C. DOCUMENTS CONSIDERED TO BE RELEVANT Relevant to claim No. Citation of document, with indication, where appropriate, of the relevant passages WO 96 30389 A (MILLENIUM PHARMACEUTICALS, 1-60 INC.; SHYJAN A.) 3 October 1996 see page 112 - page 127 WO 96 02552 A (CYTOCLONYL PHARMACEUTICS, 1-60 INC.; TORCZYNSKI R. ET AL.) 1 February 1996 see the whole document 1,2,4-7 YOU L ET AL.: "Identification of early growth response gene-1 (Egr-1) as a phorbol myristate-induced gene in lung cancer cells by differential mRNA display" AM. J. RESPIR. CELL MOL. BIOL., vol. 17, no. 5, November 1997, pages 617-624, XP002106654 see page 618, left-hand column, paragraph -/--Patent family members are listed in annex. X Further documents are listed in the continuation of box C. Special categories of cited documents: ater document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the "A" document defining the general state of the art which is not considered to be of particular relevance earlier document but published on or after the international filing date "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alor "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) document of particular relevance; the claimed invention cannot be considered to involve an inventive: step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art. Of document referring to an oral disclosure, use, exhibition or document published prior to the international filing date but later than the priority date claimed *&* document member of the same patent family Date of mailing of the international search report Date of the actual completion of the international search 22 10 1999 21 June 1999 **Authorized officer** Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentiaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo ni, CUPIDO, M Fax: (+31-70) 340-3016

	Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)	
This Inte	rmational Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:	-
1. X	Claims Nos.:	
	because they relate to subject matter not required to be searched by this Authority, namely:	
	Remark: Although claims 16, 17, 24-26, 32, 33, 48-53 and 56-58 are directed to a method of treatment of the human/animal body the search has been carried out and based on the alleged effects of the composition.	·
2. 🔲	Claims Nos.:	. :
	because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:	
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3.	Claims Nos.:	
. —	because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).	
Box II	Observations where unity of invention is lacking (Continuation of item 2 of first sheet)	
This Inte	ernational Searching Authority found multiple inventions in this international application, as follows:	
se	e FURTHER INFORMATION sheet	_
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" —	As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.	
2. 🔲	As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment	· £
	of any additional fee.	•
s. 🗀	As only some of the required additional search fees were timely paid by the applicant, this International Search Report	•
L	covers only those claims for which fees were paid, specifically claims Nos.:	
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ك ٠	No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:	
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	see FURTHER INFORMATION sheet, subject 1.	•
Barrel	on Protest The additional search fees were accompanied by the applicant's protest.	
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INTERNATIONAL SEARCH REPORT

information on patent family members

tnts 'onal Application No
PCT/US 99/01642

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